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#### (57) Abstract

A method for identifying suitable targets for antibacterial agents based on identifying targets of bacteriophage-encoded proteins is described. Also described are compositions useful in the identification methods and in inhibiting bacterial growth, and methods for preparing and using such compositions.

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# DESCRIPTION

# Development of Novel Anti-Microbial Agents Based on Bacteriophage Genomics

# BACKGROUND OF THE INVENTION

The present invention relates to the field of antibacterial agents and the treatment of infections of animals or other complex organisms by bacteria.

The frequency and spectrum of antibiotic-resistant infections have, in recent years, increased in both the hospital and community. Certain infections have become essentially untreatable and are growing to epidemic proportions in the developing world as well as in institutional settings in the developed world. The staggering spread of antibiotic resistance in pathogenic bacteria has been attributed to microbial genetic characteristics, widespread use of antibiotic drugs, and changes in society that enhance the transmission of drug-resistant organisms. This spread of drug resistant microbes is leading to ever increasing morbidity, mortality and health-care costs.

Ironically, it is the very success of antibiotics, resulting in their widespread use, that has contributed the most to rising numbers of drug resistant bacterial strains. The longer a bacterial strain is exposed to a drug, the more likely it is to acquire resistance. Today, a total of 160 antibiotics, all based on a few basic chemical structures and targeting a small number of metabolic pathways, have found their way to market. Over-prescription of these drugs, as well as the failure of patients to comply with the complete antibiotic regimen, has lead to the rapid emergence of antibiotic resistant strains. Such misuse of prescriptions, careless use of antibiotics in virtually all commercial production of beef and fowl, and changing societal conditions, such as the growth of day-care centers, increased long-term care in hospitals, and increased mobility of the population, has provided an environment where drug-resistant microbes can emerge and spread. Thus, virtually all common infectious bacteria are becoming, or have already become, resistant to one or more groups of antibiotics. Such resistance now reaches all classes of antibiotics currently in use, including: β-lactams, fluoroquinolones, aminoglycosides, macrolide peptides, chloramphenicol, tetracyclines, rifampicin, folate inhibitors, glycopeptides, and mupirocin.

Over the last 45 years bacteria have adapted genetically to avoid the destruction/alteration of the essential pathways that these chemotherapeutic agents

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target. Antibiotic resistant bacterial strains are now emerging at a higher rate than the rate at which new antibiotics are being developed. The consequence of this dilemma has been a dramatic increase in the cost of treating infections what would otherwise easily succumb to routine antibiotic therapy. Furthermore, and perhaps most importantly, the emergence of multiple drug resistant pathogenic bacteria has led to a significant increase in morbidity and mortality, particularly in institutional settings.

Most major pharmaceutical companies have on-going drug discovery programs for novel anti-microbials. These are based on screens for small molecule inhibitors (natural products, bacterial culture media, libraries of small molecules, combinatorial chemistry) of crucial metabolic pathways of the micro-organism of interest (e.g., bacteria, fungi, parasites, worms). The screening process is largely for cytotoxic compounds and in most cases is not based on a known mechanism of action of the compounds. Pharmaceutical companies have large programs in this area. Classical drug screening programs are being exhausted and many of these pharmaceutical companies are looking towards rational drug design programs.

Several small to mid-size biotechnology companies as well as large pharmaceutical companies have developed systematic high-throughput sequencing programs to decipher the genetic code of specific micro-organisms of interest. The goal is to identify, through sequencing, unique biochemical pathways or intermediates that are unique to the microorganism. Knowledge of this may, in turn, form the rationale for a drug discovery program based on the mechanism of action of the identified enzymes/proteins. Genome Therapeutics Corp., The Institute for Genome Research, Human Genome Sciences Inc., and other companies have such sequencing programs in place. However, one of the most critical steps in this approach is the ascertainment that the identified proteins and biochemical pathways are 1) non-redundant and essential for bacterial survival, and 2) constitute suitable and accessible targets for drug discovery.

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# SUMMARY OF THE INVENTION

While animals such as humans are, on occasion, infected by pathogenic bacteria, bacteria also have natural enemies. A number of host-specific viruses, known as bacteriophages or phages, infect and kill bacteria in the natural environment. Such bacteriophages generally have small compact genomes and bacteria are their exclusive hosts. Many known bacteria are host to a large number of bacteriophages that have been described in the literature. During the 1940's - 1960's, phage biology was an area of active research. As a testimony to this, the study of phages which infect and inhibit the enteric bacterium *Escherichia coli* (*E. coli*) contributed much to the early understanding of molecular biology and virology.

As is generally understood, bacteriophage (or phages) are viruses that infect and kill bacteria. They are natural enemies of bacteria and, over the course of evolution, have developed proteins (products of DNA sequences) which enable them to infect a host bacteria, replicate their genetic material, usurp host metabolism, and ultimately kill their host. The scientific literature well documents the fact that many known bacteria have a large number of such bacteriophages (Ackermann and DuBow, 1987) that can infect and kill them (for example, see the ATCC bacteriophage collection at http://www.atcc.org).

This invention utilizes the observation that bacteriophages successfully infect and inhibit or kill host bacteria, targeting a variety of normal host metabolic and physiological traits, some of which are shared by all bacteria, pathogenic and nonpathogenic alike. The term "pathogenic" as used herein denotes a contribution to or implication in disease or a morbid state of an infected organism. The invention thus involves identifying and elucidating the molecular mechanisms by which phages interfere with host bacterial metabolism, an objective being to provide novel targets for drug design. Whether the phage blocks bacterial RNA transcription or translation, or attacks other important metabolic pathways, such as cell wall assembly or membrane integrity, the basic blueprint for a phage's bacteria-inhibiting ability is encoded in its genome and can be unlocked using bioinformatics, functional genomics, and proteomics. By these means, the invention utilizes sequence information from the genomics of bacteriophage to identify novel antimicrobials that can be further used to actively and/or prophylactically treat bacterial infection.

Two important components of the invention thus are: i) the identification of bacteria-inhibiting phage open reading frames ("ORF"s) and corresponding products that can be used to develop antibiotics based on amino acid sequence and secondary structural characteristics of the ORF products, and ii) the use of bacteriophages to map

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out essential bacterial target genes and homologs, which can in turn lead to the development of suitable anti-microbial agents. These two avenues represent new and general methods for developing novel antimicrobials.

The invention thus concerns the identification of bacteriophage ORFs that supply bacteria-inhibiting functions. In this regard, use of the terms "inhibit", "inhibition", "inhibitory", and "inhibitor" all refer to a function of reducing a biological activity or function. Such reduction in activity or function can, for example, be in connection with a cellular component, e.g., an enzyme, or in connection with an overall process, e.g., synthesis of a particular protein, or in connection with an overall process of a cell, e.g., cell growth. In reference to bacterial cell growth, for example, an inhibitory effect (i.e., a bacteria-inhibiting effect) may be bacteriocidal (killing of bacterial cells) or bacteriostatic (i.e., stopping or at least slowing bacterial cell growth). The latter slows or prevents cell growth such that fewer cells of the strain are produced relative to uninhibited cells over a given period of time. From a molecular standpoint, such inhibition may equate with a reduction in the level of, or elimination of, the transcription and/or translation of a specific bacterial target(s), or reduction or elimination of activity of a particular target biomolecule.

It is particularly advantageous to evaluate a plurality of different phage ORFs for inhibitory activity that may be from one, but is preferably from a plurality of different phage. For example, evaluating ORFs from a number of different phage of the same bacterial host provides at least two advantages. One is that the multiple phages will provide identification of a variety of different targets. Second, it is likely that multiple phage will utilize the same cellular target

As used herein, the terms "bacteriophage" and "phage" are used interchangeably to refer to a virus which can infect a bacterial strain or a number of different bacterial strains.

In the context of this invention, the term "bacteriophage ORF" or "phage ORF" or similar term refers to a nucleotide sequence in or from a bacteriophage. In connection with a particular ORF, the terms refer an open reading frame which has at least 95% sequence identity, preferably at least 97% sequence identity, more preferably at least 98% sequence identity with an ORF from the particular phage identified herein (e.g., with an ORF as identified herein) or to a nucleic acid sequence which has the specified sequence identify percentage with such an ORF sequence.

A first aspect of the invention thus provides a method for identifying a bacteriophage nucleic acid coding region encoding a product active on an essential bacterial target by identifying a nucleic acid sequence encoding a gene product which

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provides a bacteria-inhibiting function when the bacteriophage infects a host bacterium, preferably one that is an animal or plant pathogen, more preferably a bird or mammalian pathogen, and most preferably a human pathogen. The bacteriophage is an uncharacterized bacteriophage. Thus, the method excludes, for example, phage  $\lambda$ ,  $\phi$ x174, m13 and other *E.coli*-specific bacteriophage that have been studied with respect to gene number and/or function. It also excludes, for example, the nucleic

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acid coding regions described in Tables 12-14, and in preferred embodiments, excludes the phage in which those regions are naturally located.

In connection with bacteriophage, the term "uncharacterized" means that a certain bacteriophage's genome has not yet been fully identified such that the genes having function involved in inhibiting host cells have not been identified. In particular, phage for which the description of genomic or protein sequence was first provided herein are uncharacterized. Phage sequences for which host bacteriainhibiting functions have been identified prior to the filing of the present application (or alternatively prior to the present invention) are specifically excluded from the aspects involving utilization of sequences from uncharacterized bacteriophage, except that aspects may involve a plurality of phage where one or more of those phage are uncharacterized and one or more others have been characterized to some extent. A number of different bacteria-inhibiting phage ORFs are indicated in Tables 11-14. The phage ORFs or sequences identified therein are not within the term "uncharacterized; alternatively, in preferred embodiments the phage containing those ORFs are excluded from this term. Further, any additional phage ORFs (or alternatively the phage which contain those ORFs) which have previously been described in the art as bacteria-inhibiting ORFs are expressly excluded; those ORFs or phage are known to those skilled in the art and the exclusion can be made express by specifically naming such ORFs or phage as needed (likewise for uncharacterized targets as described below). For the sake of brevity, such a listing is not expressly presented, as such information is readily available to those skilled in the art.

Stating that an agent or compound is "active on" a particular cellular target, such as the product of a particular gene, means that the target is an important part of a cellular pathway which includes that target and that the agent acts on that pathway. Thus, in some cases the agent may act on a component upstream or downstream of the stated target, including on a regulator of that pathway or a component of that pathway.

By "essential", in connection with a gene or gene product, is meant that the host cannot survive without, or is significantly growth compromised, in the absence depletion, or alteration of functional product. An "essential gene" is thus one that encodes a product that is beneficial, or preferably necessary, for cellular growth in

vitro in a medium appropriate for growth of a strain having a wild-type allele corresponding to the particular gene in question. Therefore, if an essential gene is inactivated or inhibited, that cell will grow significantly more slowly, preferably less than 20%, more preferably less than 10%, most preferably less than 5% of the growth rate of the uninhibited wild-type, or not at all, in the growth medium. Preferably, in the absence of activity provided by a product of the gene, the cell will not grow at all or will be non-viable, at least under culture conditions similar to the *in vivo* conditions normally encountered by the bacterial cell during an infection. For example, absence of the biological activity of certain enzymes involved in bacterial cell wall synthesis can result in the lysis of cells under normal osmotic conditions, even though protoplasts can be maintained under controlled osmotic conditions. In the context of the invention, essential genes are generally the preferred targets of antimicrobial agents. Essential genes can encode target molecules directly or can encode a product involved in the production, modification, or maintenance of a target molecule.

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A "target" refers to a biomolecule that can be acted on by an exogenous agent, thereby modulating, preferably inhibiting, growth or viability of a cell. In most cases such a target will be a nucleic acid sequence or molecule, or a polypeptide or protein. However, other types of biomolecules can also be targets, e.g., membrane lipids and cell wall structural components.

The term "bacterium" refers to a single bacterial strain, and includes a single cell, and a plurality or population of cells of that strain unless clearly indicated to the contrary. In reference to bacteria or bacteriophage, the term "strain" refers to bacteria or phage having a particular genetic content. The genetic content includes genomic content as well as recombinant vectors. Thus, for example, two otherwise identical bacterial cells would represent different strains if each contained a vector, e.g., a plasmid, with different phage ORF inserts.

In preferred embodiments, the phage is *Staphylococcus aureus* phage 77, 3A, 96, or 44 AHJD, *Enterococcus* sp. phage 182, or *Streptococcus pneumoniae* phage Dp-1.

In preferred embodiments, the phage is selected from. Preferred embodiments involve expressing at least one recombinant phage ORF(s) in a bacterial host followed by inhibition analysis of that host. Inhibition following expression of the phage ORF is indicative that the product of the ORF is active on an essential bacterial target. Such evaluation can be carried out in a variety of different formats, such as on a support matrix such as a solidified medium in a petri dish, or in liquid culture.

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Preferably a plurality of phage ORFs are expressed in at least one bacterium. The plurality of phage ORFs can be from one or a plurality of phage. With respect to a single phage or at least one phage in a plurality of phages, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome. Preferably, for a plurality of phage, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome of each phage. The plurality of phage ORFs can be expressed in a single bacterium, or in a plurality of bacteria where one ORF is expressed in each bacterium, or in a plurality of bacteria where a plurality of ORFs are expressed in at least one or in all of the plurality of bacteria, or combinations of these.

In embodiments of the above aspect (as well as in other aspects herein) in which a plurality of phage are utilized, a plurality of phage have the same bacterial host species; have different bacterial host species; or both. The plurality of phage includes at least two different phage, preferably at least 3,4,5,6,8,10,15,20, or more different phage. Indeed, more preferably, the plurality of phage will include 50, 75, 100, or more phage. As described herein, the larger number of phage is useful to provide additional target and target evaluation information useful in developing antibacterial agents, for example, by providing identification of a larger range of bacterial targets, and/or providing further indication of the suitability of a particular target (for example, utilization of a target by a number of different unrelated phage can suggest that the target is particularly stable and accessible and effective) and/or can indicate alternate sites on a target which interact with different inhibitors.

Further embodiments involve confirmation of the inhibitor function of the phage ORF, such as by utilizing or incorporating a control(s) designed to confirm the inhibitory nature of the ORF(s) being evaluated. The control can, for example, be provided by expression of an inactive or partially inactive form of the ORF or ORF product, and/or by the absence of expression of the ORF or ORF product in the same or a closely comparable bacterial strain as that used for expression of the test ORF. The reduced level of activity or the absence of active ORF product in the control will thus not provide the inhibition provided by a corresponding inhibitory ORF, or will provide a distinguishably lower level of inhibition. An inactivated or partially inactivated control has a mutation(s), e.g., in the coding region or in flanking regulatory elements, that reduce(s) or eliminate(s) the normal function of the ORF.

Thus, the inhibition of a bacterium following expression of a phage ORF is determined by comparison with the effects of expression of an inactivated ORF or the

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response of the bacteria in the absence of expression in the same or similar type bacterium. Such determination of inhibition of the bacterium following expression of the ORF is indicative of a bacteria-inhibiting function. These manipulations are routinely understood and accomplished by those of skill in the art using standard techniques. In embodiments utilizing absence of expression of the ORF, the bacteria can, for example, contain an empty vector or a vector which allows expression of an unrelated sequence which is preferably non-inhibitory. Alternatively, the bacteria may have no vector at all. Combinations of such controls or other controls may also be utilized as recognized by those skilled in the art.

In embodiments involving expression of a phage ORF in a bacterial strain, in preferred embodiments that expression is inducible.

By "inducible" is meant that expression is absent or occurs at a low level until the occurrence of an appropriate environmental stimulus provides otherwise. For the present invention such induction is preferably controlled by an artificial environmental change, such as by contacting a bacterial strain population with an inducing compound (i.e., an inducer). However, induction could also occur, for example, in response to build-up of a compound produced by the bacteria in the bacterial culture, e.g., in the medium. As uncontrolled or constitutive expression of inhibitory ORFs can severely compromise bacteria to the point of eradication, such expression is therefore undesirable in many cases because it would prevent effective evaluation of the strain and inhibitor being studied. For example, such uncontrolled expression could prevent any growth of the strain following insertion of a recombinant ORF, thus preventing determination of effective transfection or transformation. A controlled or inducible expression is therefore advantageous and is generally provided through the provision of suitable regulatory elements, e.g., promoter/operator sequences that can be conveniently transcriptionally linked to a coding sequence to be evaluated. In most cases, the vector will also contain sequences suitable for efficient replication of the vector in the same or different host cells and/or sequences allowing selection of cells containing the vector, i.e., "selectable markers." Further, preferred vectors include convenient primer sequences flanking the cloning region from which PCR and/or sequencing may be performed.

As knowledge of the nucleotide sequence of phage ORFs is useful, e.g., for assisting in the identification of phage proteins active against essential bacterial host targets, preferred embodiments involve the sequencing of at least a portion of the phage genome in combination with the above methods. This can be done either before or after or independent of expression and inhibition of the ORF in the bacteria, and provides information on the nature and characteristics of the ORF. Such a portion is

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preferably at least 10%, 20%, 40%, 80%, 90%; or 100% of the phage genome. For embodiments in which a plurality of phage are utilized, preferably each phage is sequenced to an extent as just specified.

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Such sequencing is preferably accompanied by computer sequence analysis to define and evaluate ORF(s), ORF products, structural motifs or functional properties of ORF products, and/or their genetic control elements. Thus, certain embodiments incorporate computer sequence analyses or nucleic acid and/or amino acid sequences. Further, existing data banks can provide phage sequence and product information which can be utilized for analysis and identification of ORFs in the sequence. Computer analysis may further employ known homologous sequences from other species that suggest or indicate conserved underlying biochemical function(s) for the inhibitory or potentially inhibitory ORF sequence(s) being evaluated. This can include the sequences of signature motifs of identified classes of inhibitors.

In the context of the phage nucleic acid sequences, e.g., gene sequences, of this invention, the terms "homolog" and "homologous" denote nucleotide sequences from different bacteria or phage strains or species or from other types of organisms that have significantly related nucleotide sequences, and consequently significantly related encoded gene products, preferably having related function. Homologous gene sequences or coding sequences have at least 70% sequence identity (as defined by the maximal base match in a computer-generated alignment of two or more nucleic acid sequences) over at least one sequence window of 48 nucleotides, more preferably at least 80 or 85%, still more preferably at least 90%, and most preferably at least 95%. The polypeptide products of homologous genes have at least 35% amino acid sequence identity over at least one sequence window of 18 amino acid residues, more preferably at least 40%, still more preferably at least 50% or 60%, and most preferably at least 70%, 80%, or 90%. Preferably, the homologous gene product is also a functional homolog, meaning that the homolog will functionally complement one or more biological activities of the product being compared. For nucleotide or amino acid sequence comparisons where a homology is defined by a % sequence identity, the percentage is determined using BLAST programs ( with default parameters (Altschul et al., 1997, "Gapped BLAST and PSI-BLAST: a new generation of protein database search programs, Nucleic Acid Res. 25:3389-3402). Any of a variety of algorithms known in the art which provide comparable results can also be used, preferably using default parameters. Performance characteristics for three different algorithms in homology searching is described in Salamov et al., 1999, "Combining sensitive database searches with multiple intermediates to detect distant

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homologues." *Protein Eng.* 12:95-100. Another exemplary program package is the GCG<sup>TM</sup> package from the University of Wisconsin.

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Homologs may also or in addition be characterized by the ability of two complementary nucleic acid strands to hybridize to each other under appropriately stringent conditions. Hybridizations are typically and preferably conducted with probe-length nucleic acid molecules, preferably 20-100 nucleotides in length. Those skilled in the art understand how to estimate and adjust the stringency of hybridization conditions such that sequences having at least a desired level of complementarity will stably hybridize, while those having lower complementarity will not. For examples of hybridization conditions and parameters, see, e.g., Maniatis, T. et al. (1989)

Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology.

John Wiley & Sons, Secaucus, N.J. Homologs and homologous gene sequences may thus be identified using any nucleic acid sequence of interest, including the phage ORFs and bacterial target genes of the present invention.

A typical hybridization, for example, utilizes, besides the labeled probe of interest, a salt solution such as 6xSSC (NaCl and Sodium Citrate base) to stabilize nucleic acid strand interaction, a mild detergent such as 0.5% SDS, together with other typical additives such as Denhardt's solution and salmon sperm DNA. The solution is added to the immobilized sequence to be probed and incubated at suitable temperatures to preferably permit specific binding while minimizing nonspecific binding. The temperature of the incubations and ensuing washes is critical to the success and clarity of the hybridization. Stringent conditions employ relatively higher temperatures, lower salt concentrations, and/or more detergent than do non-stringent conditions. Hybridization temperatures also depend on the length, complementarity level, and nature (ie, "GC content") of the sequences to be tested. Typical stringent hybridizations and washes are conducted at temperatures of at least 40°C, while lower stringency hybridizations and washes are typically conducted at 37°C down to room temperature (~25°C). One of skill in the art is aware that these conditions may vary according to the parameters indicated above, and that certain additives such as formamide and dextran sulphate may also be added to affect the conditions.

By "stringent hybridization conditions" is meant hybridization conditions at least as stringent as the following: hybridization in 50% formamide, 5X SSC, 50 mM NaH,PO,, pH 6.8, 0.5% SDS, 0.1 mg/mL sonicated salmon sperm DNA, and 5X Denhart's solution at 42°C overnight; washing with 2X SSC, 0.1% SDS at 45°G; and washing with 0.2X SSC, 0.1% SDS at 45°C.

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In sequence comparison analyses, an ORF, or motif, or set of motifs in a bacteriophage sequence can be compared to known inhibitor sequences, e.g., homologous sequences encoding homologous inhibitors of bacterial function. Likewise, the analysis can include comparison with the structure of essential bacterial gene products, as structural similarities can be indicative of similar or replacement biological function. Such analysis can include the identification of a signature, or characteristic motif(s) of an inhibitor or inhibitor class.

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Also, the identification of structural motifs in an encoded product, based on nucleotide or amino acid sequence analysis, can be used to infer a biochemical function for the product. A database containing identified structural motifs in a large number of sequences is available for identification of motifs in phage sequences. The database is PROSITE, which is available at www.expasy.ch/cgi~bin/scanprosite. The identification of motifs can, for example, include the identification of signature motifs for a class or classes of inhibitory proteins. Other such databases may also be used.

In aspects and preferred embodiments described herein, in which a bacterium or host bacterium is specified, the bacterium or host bacterium is preferably selected from a pathogenic bacterial species, for example, one selected from Table 1. Preferably, an animal or plant pathogen is used. For animals, preferably the bacterium is a bird or mammalian pathogen, still more preferably a human pathogen.

In aspects and preferred embodiments involving a bacteriophage or sequences from a bacteriophage, one or more bacteriophage are preferably selected from those listed in Table 1. Those exemplary bacteriophage are readily obtained from the indicated sources.

In some cases, it is advantageous to utilize phage with non-pathogenic host bacteria. The genome, structural motif, ORF, homolog, and other analyses described herein can be performed on such phage and bacteria. Such analysis provides useful information and compositions. The results of such analyses can also be utilized in aspects of the present invention to identify homologous ORFs, especially inhibitor ORFs in phage with pathogenic bacterial hosts. Similarly, identification of a target in a non-pathogenic host can be used to identify homologous sequences and targets in pathogenic bacteria, especially in genetically closely related bacteria. Those skilled in the art are familiar with bacterial genetic relationships and with how to determine relatedness based on levels of genomic identity or other measures of nucleotide sequence and/or amino acid sequence similarity, and/or other physical and culture characteristics such as morphology, nutritional requirements, or minimal media-to support growth.

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Also in preferred embodiments, an embodiments of this aspect is combined with an embodiment of the following aspect.

A related aspect of the invention provides methods for identifying a target for antibacterial agents by identifying the bacterial target(s) of at least one uncharacterized or untargeted inhibitor protein or RNA from a bacteriophage. Such identification allows the development of antibacterial agents active on such targets. Preferred embodiments for identifying such targets involve the identification of binding of target and phage ORF products to one another. The phage ORF products may be subportions of a larger ORF product that also binds the host target. In preferred embodiments, the phage protein or RNA is from an uncharacterized bacteriophage in Table 1. This aspect preferably includes the identification of a plurality of such targets in one or a plurality of different bacteria, preferably in one or a plurality of bacteria listed in Table 1.

In preferred embodiments of this aspect and other aspects of this invention involving particular phage ORFs or phage sequences, the ORF is *Staphylococcus aureus* phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As indicated for the above aspect, preferably the method involves the use of a plurality of different phage, and thus a plurality of different phage inhibitors and/or inhibitor ORFs.

In addition to uncharacteized phage ORF products, it is also useful to identify the targets of phage ORF products which are known to be inhibitors of host bacteria, but where the target has not been identified. Thus, such inhibitors can likewise be utilized as "untargeted" inhibitor phage ORFs and ORF products, e.g., proteins or RNAs.

In the context of inhibitor proteins or RNAs from a phage, the term "uncharacterized" means that a bacteria-inhibiting function for the protein has not previously been identified. Preferably, but not necessarily, the sequence of the protein or the corresponding coding region or ORF was not described in the art before the filing of the present application for patent (or alternatively prior to the present invention). Thus, this term specifically excludes any bacteria-inhibiting phage protein and its associated bacterial target which has been identified as inhibitory before the present invention or alternatively before the filing of the present application, for example those identified in Tables 12-14 or otherwise identified herein. For example, from *E. coli*, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, phage T4

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gp55/gp33 alter the specificity of host RNA polymerase. The T4 regB gene product also targets the host translation apparatus. As with the uncharacterized bacteriophage ORFs or bacteriophage above, for such identified proteins, the sequences encoding those proteins are excluded from the uncharacterized inhibitor proteins.

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The term "fragment" refers to a portion of a larger molecule or assembly. For proteins, the term "fragment" refers to a molecule which includes at least 5 contiguous amino acids from the reference polypeptide or protein, preferably at least 8, 10, 12, 15, 20, 30, 50 or more contiguous amino acids. In connection with oligo- or polynucleotides, the term "fragment" refers to a molecule which includes at least 15 contiguous nucleotides from a reference polynucleotide, preferably at least 24, 30, 36, 45, 60, 90, 150, or more contiguous nucleotides.

Preferred embodiments involve identification of binding that include methods for distinguishing bound molecules, for example, affinity chromatography, immunoprecipitation, crosslinking, and/or genetic screen methods that permit protein:protein interactions to be monitored. One of skill in the art is familiar with these techniques and common materials utilized (see, e.g., Coligan, J. et al. (eds.) (1995) Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J.).

Genetic screening for the identification of protein:protein interactions typically involves the co-introduction of both a chimeric bait nucleic acid sequence (here, the phage ORF to be tested) and a chimeric target nucleic acid sequence that, when co-expressed and having affinity for one another in a host cell, stimulate reporter gene expression to indicate the relationship. A "positive" can thus suggest a potential inhibitory effect in bacteria. This is discussed in further detail in the Detailed Description section below. In this way, new bacterial targets can be identified that are inhibited by specific phage ORF products or derivatives, fragments, mimetics, or other molecules.

Other embodiments involve the identification and/or utilization of mutant targets by virtue of their host's relatively unresponsive nature in the presence of expression of ORFs previously identified as inhibitory to the non-mutant or wild-type strain. Such mutants have the effect of protecting the host from an inhibition that would otherwise occur and indirectly allow identification of the precise responsible target for follow-up studies and anti-microbial development. In certain embodiments, rescue from inhibition occurs under conditions in which a bacterial target or mutant target is highly expressed. This is performed, for example, through coupling of the sequence with regulatory element promoters, e.g., as known in the art, which regulate expression at levels higher than wild-type, e.g., at a level sufficiently higher that the

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inhibitor can be competitively bound to the highly expressed target such that the bacterium is detectably less inhibited.

Identification of the bacterial target can involve identification of a phage-specific site of action. This can involve a newly identified target, or a target where the phage site of action differs from the site of action of a previously known antibacterial agent or inhibitor. For example, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, which is also the cellular target for the antibacterial agent, rifampin. To the extent that a phage product is found to act at a different site than previously described inhibitors, aspects of the present invention can utilize those new, phage-specific sites for identification and use of new agents. The site of action can be identified by techniques well-known to those skilled in the art, for example, by mutational analysis, binding competition analysis, and/or other appropriate techniques.

Once a bacterial host target protein or nucleic acid or mutant target sequence has been identified and/or isolated, it too can be conveniently sequenced, sequence analyzed (e.g., by computer), and the underlying gene(s), and corresponding translated product(s) further characterized. Preferred embodiments include such analysis and identification. Preferably such a target has not previously been identified as an appropriate target for antibacterial action.

Certain embodiments include the identification of at least one inhibitory phage ORF or ORF product, e.g., as described for the above aspect, and thus are a combination of the two aspects.

Additionally, the invention provides methods for identifying targets for antibacterial agents by identifying homologs of a bacterial target e.g., S. aureus, Enterococcus faecalis or other Enterococci, and Streptococcus pneumoniae of a bacteriophage inhibitory ORF product. Such homologs may be utilized in the various aspects and embodiments described herein as described for the host Enterococcus sp. for bacteriophage 182.

Other aspects of the invention provide isolated, purified, or enriched specific phage nucleic acid and amino acid sequences, subsequences, and homologs thereof for phage selected from uncharacterized phage listed in Table 1, preferably from bacteriophage 77, 3A, 96, 44AHJD (Staphylococcus aureus host bacterium), Dp-1 (Streptococcus pneumoniae host), or 182 (Enterococcus host) or other phage listed in Table 1 for those bacteria. For example, such sequences do not include sequences identified in any of Tables 11-14. Nucleotide sequences of this aspect are at least 15 nucleotides in length, preferably at least 18, 21, 24, or 27 nucleotides in length, more preferably at least 30, 50, or 90 nucleotides in length. In certain embodiments, longer

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nucleic acids are preferred, for example those of at least 120, 150, 200, 300, 600, 900 or more nucleotides. Such sequences can, for example, be amplification oligonucleotides (e.g., PCR primers), oligonucleotide probes, sequences encoding a portion or all of a phage-encoded protein, or a fragment or all of a phage-encoded protein. In preferred embodiments, the nucleic acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF. The upper length limit can also be expressed in terms of the number of base pairs of the ORF (coding region). In preferred embodiments, the nucleic acid sequence is from Staphylococcus aureus phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, S. aureus phage 44 AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

As it is recognized that alternate codons will encode the same amino acid for

most amino acids due to the degeneracy of the genetic code, the sequences of this aspect includes nucleic acid sequences utilizing such alternate codon usage for one or more codons of a coding sequence. For example, all four nucleic acid sequences GCT, GCC, GCA, and GCG encode the amino acid, alanine. Therefore, if for an amino acid there exists an average of three codons, a polypeptide of 100 amino acids in length will, on average, be encoded by  $3^{100}$ , or 5 x  $10^{47}$ , nucleic acid sequences. Thus, a nucleic acid sequence can be modified (e.g., a nucleic acid sequence from a phage as specified above) to form a second nucleic acid sequence encoding the same polypeptide as encoded by the first nucleic acid sequence using routine procedures and without undue experimentation. Thus, all possible nucleic acid sequences that encode the specified amino acid sequences are also fully described herein, as if all were written out in full, taking into account the codon usage, especially that preferred in the host bacterium. The alternate codon descriptions are available in common texbooks, for example, Stryer, BIOCHEMISTRY 3rd ed., and Lehninger, BIOCHEMISTRY 3rd ed., along wth many others. Codon preference tables for various types of organisms are available in the literature. Sequences with alternate codons at one or more sites can also be utilized in the computer-related aspects and embodiments herein. Because of the number of sequence variations involving alternate codon usage, for the sake of brevity, individual sequences are not separately listed herein. Instead the alternate sequences are described by reference to the natural sequence with replacement of one or more (up to all e.g., up to 3, 5, 10, 15, 20, 30, 40, 50, or more) of the degenerate codons with alternate codons from the alternate codon

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table (Table 6), or a modified table applicable to a particular organism that has differing codon usage, preferably with selection according to preferred codon usage for the normal host organism or a host organism in which a sequence is intended to be expressed. Those skilled in the art also understand how to alter the alternate codons to be used for expression in organisms where certain codons code differently than shown in the "universal" codon table.

For amino acid sequences or polypeptides, sequences contain at least 5 peptide-linked amino acid residues, and preferably at least 6, 7, 10, 15, 20, 30, or 40, amino acids having identical amino acid sequence as the same number of contiguous amino acid residues in a particular phage ORF product. In some cases longer sequences may be preferred, for example, those of at least 50, 60, 70, 80, or 100 amino acids in length. In preferred embodiments, the amino acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF product. The upper length limit can also be expressed in terms of the number of amino acid residues of the ORF product. In preferred embodiments, the amino acid sequence or polypeptide has bacteria-inhibiting function when expressed or otherwise present in a bacterial cell which is a host for the bacteriophage from which the sequence was derived.

By "isolated" in reference to a nucleic acid is meant that a naturally occurring sequence has been removed from its normal cellular (e.g., chromosomal) environment or is synthesized in a non-natural environment (e.g., artificially synthesized). Thus, the sequence may be in a cell-free solution or placed in a different cellular environment. The term does not imply that the sequence is the only nucleotide chain present, but that it is essentially free (about 90-95% pure at least) of non-nucleotide material naturally associated with it, and thus is distinguished from isolated chromosomes.

The term "enriched" means that the specific DNA or RNA sequence constitutes a significantly higher fraction (2-5 fold) of the total DNA or RNA present in the cells or solution of interest than in normal or diseased cells or in cells from which the sequence was originally taken. This could be caused by a person by preferential reduction in the amount of other DNA or RNA present, or by a preferential increase in the amount of the specific DNA or RNA sequence, or by a combination of the two. However, it should be noted that enriched does not imply that there are no other DNA or RNA sequences present, just that the relative amount of the sequence of interest has been significantly increased.

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The term "significant" is used to indicate that the level of increase is useful to the person making such an increase and an increase relative to other nucleic acids of about at least 2-fold, more preferably at least 5- to 10-fold or even more. The term also does not imply that there is no DNA or RNA from other sources. The other source DNA may, for example, comprise DNA from a yeast or bacterial genome, or a cloning vector such as pUC19. This term distinguishes from naturally occurring events, such as viral infection, or tumor type growths, in which the level of one mRNA may be naturally increased relative to other species of mRNA. That is, the term is meant to cover only those situations in which a person has intervened to elevate the proportion of the desired nucleic acid.

It is also advantageous for some purposes that a nucleotide sequence be in purified form. The term "purified" in reference to nucleic acid does not require absolute purity (such as a homogeneous preparation). Instead, it represents an indication that the sequence is relatively more pure than in the natural environment (compared to the natural level, this level should be at least 2-5 fold greater, e.g., in terms of mg/mL). Individual clones isolated from a cDNA library may be purified to electrophoretic homogeneity. The claimed DNA molecules obtained from these clones could be obtained directly from total DNA or from total RNA. The cDNA clones are not naturally occurring, but rather are preferably obtained via manipulation of a partially purified naturally occurring substance (messenger RNA). The construction of a cDNA library from mRNA involves the creation of a synthetic substance (cDNA) and pure individual cDNA clones can be isolated from the synthetic library by clonal selection of the cells carrying the cDNA library. Thus, the process which includes the construction of a cDNA library from mRNA and isolation of distinct cDNA clones yields an approximately 106-fold purification of the native message. Thus, purification of at least one order of magnitude, preferably two or three orders, and more preferably four or five orders of magnitude is expressly contemplated.

The terms "isolated", "enriched", and "purified" as respect nucleic acids, above, may similarly be used to denote the relative purity and abundance of polypeptides (multimers of amino acids joined one to another by α-carboxyl:α-amino group (peptide) bonds). These, too, may be stored in, grown in, screened in, and selected from libraries using biochemical techniques familiar in the art. Such polypeptides may be natural, synthetic or chimeric and may be extracted using any of a variety of methods, such as antibody immunoprecipitation, other "tagging" — techniques, conventional chromatography and/or electrophoretic methods. Some of the above utilize the corresponding nucleic acid sequence.

As indicated above, aspects and embodiments of the invention are not limited to entire genes and proteins. The invention also provides and utilizes fragments and portions thereof, preferably those which are "active" in the inhibitory sense described above. Such peptides or oligopeptides and oligo or polynucleotides have preferred lengths as specified above for nucleic acid and amino acid sequences from phage; corresponding recombinant constructs can be made to express the encoded same. Also included are homologous sequences and fragments thereof.

Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art. In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Also, by having particular phage ORFs, e.g., the phage ORFs identified herein (e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described), other antimicrobial sequences from other bacteriophage sources can be identified and isolated using methods described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

The invention also provides bacteriophage antimicrobial DNA segments from other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences that are highly homologous. The bacteriophage segment from a specific phage, e.g., an antimicrobial DNA segment, can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on nucleic acid and/or amino acid sequence comparisons. As with identified inhibitory sequences, such homologous coding sequences and products can be used as antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

The nucleotide and amino acid sequences identified herein are believed to be correct, however, certain sequences may contain a small percentage of errors, e.g., 1-5%. In the event that any of the sequences have errors, the corrected sequences can be readily provided by one skilled in the art using routine methods. For example, the nucleotide sequences can be confirmed or corrected by obtaining and culturing the relevant phage, and purifying phage genomic nucleic acids: A region or regions of interest can be amplified, e.g., by PCR from the appropriate genomic template, using primers based on the described sequence. The amplified regions can then be sequenced using any of the available methods (e.g., a dideoxy termination method).

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This can be done redundantly to provide the corrected sequence or to confirm that the described sequence is correct. Alternatively, a particular sequence or sequences can be identified and isolated as an insert or inserts in a phage genomic library and isolated, amplified, and sequenced by standard methods. Confirmation or correction of a nucleotide sequence for a phage gene provides an amino acid sequence of the encoded product by merely reading off the amino acid sequence according to the normal codon relationships and/or expressed in a standard expression system and the polypeptide product sequenced by standard techniques. The sequences described herein thus provide unique identification of the corresponding genes, coding sequences, and other sequences, allowing those sequences to be used in the various aspects of the present invention.

In other aspects, the invention provides recombinant vectors and cells harboring at least one of the phage ORFs or portion thereof, or bacterial target sequences described herein. As understood by those skilled in the art, vectors may be provided in different forms, including, for example, plasmids, cosmids, and virus-based vectors. See, e.g., Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; See also, Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J.

In preferred embodiments, the vectors will be expression vectors, preferably shuttle vectors that permit cloning, replication, and expression within bacteria. An "expression vector" is one having regulatory nucleotide sequences containing transcriptional and translational regulatory information that controls expression of the nucleotide sequence in a host cell. Preferably the vector is constructed to allow amplification from vector sequences flanking an insert locus. In certain embodiments, the expression vectors may additionally or alternativley support expression, and/or replication in animal, plant and/or yeast cells due to the presence of suitable regulatory sequences, e.g., promoters, enhancers, 3' stabilizing sequences, primer sequences, etc. In preferred embodiments, the promoters are inducible and specific for the system in which expression is desired, e.g., bacteria, animal, plant, or yeast. The vectors may optionally encode a "tag" sequence or sequences to facilitate protein purification. Convenient restriction enzyme cloning sites and suitable selective marker(s) are also optionally included. Such selective markers can be, for example, antibiotic resistance markers or markers which supply an essential nutritive growth factor to an otherwise deficient mutant host, e.g., tryptophan, histidine, or leucine in the Yeast Two-Hybrid systems described below.

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The term "recombinant vector" relates to a single- or double-stranded circular nucleic acid molecule that can be transfected into cells and replicated within or independently of a cell genome. A circular double-stranded nucleic acid molecule can be cut and thereby linearized upon treatment with appropriate restriction enzymes. An assortment of nucleic acid vectors, restriction enzymes, and the knowledge of the nucleotide sequences cut by restriction enzymes are readily available to those skilled in the art. A nucleic acid molecule encoding a desired product can be inserted into a vector by cutting the vector with restriction enzymes and ligating the two pieces together. Preferably the vector is an expression vector, e.g., a shuttle expression vector as described above.

By "recombinant cell" is meant a cell possessing introduced or engineered nucleic acid sequences, e.g., as described above. The sequence may be in the form of or part of a vector or may be integrated into the host cell genome. Preferably the cell is a bacterial cell.

In another aspect, the invention also provides methods for identifying and/or screening compounds "active on" at least one bacterial target of a bacteriophage inhibitor protein or RNA. Preferred embodiments involve contacting such a bacterial target or targets (e.g., bacterial target proteins) with a test compound, and determining whether the compound binds to or reduces the level of activity of the bacterial target (e.g., a bacterial target protein). Preferably this is done either in vivo (i.e., in a cell-based assay) or in vitro, e.g., in a cell-free system under approximately physiological conditions.

The compounds that can be used may be large or small, synthetic or natural, organic or inorganic, proteinaceous or non-proteinaceous. In preferred embodiments, the compound is a peptidomimetic, as described herein, a bacteriophage inhibitor protein or fragment or derivative thereof, preferably an "active portion", or a small molecule.

In preferred embodiments, the bacterial target is a target of a phage ORF identified herein, e.g., S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

In particular embodiments, the methods include the identification of bacterial targets or the site of action of an inhibitor on a bacterial target as described above or otherwise described herein.

In embodiments involving binding assays, preferably binding is to a fragment or portion of a bacterial target protein, where the fragment includes less than 90%, 80%, 70%, 60%, 50%, 40%, or 30% of an intact bacterial target protein. Preferably,

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the at least one bacterial target includes a plurality of different targets of bacteriophage inhibitor proteins, preferably a plurality of different targets. The plurality of targets can be in or from a plurality of different bacteria, but preferably is from a single bacterial species.

A "method of screening" refers to a method for evaluating a relevant activity or property of a large plurality of compounds (e.g., a bacteria-inhibiting activity), rather than just one or a few compounds. For example, a method of screening can be used to conveniently test at least 100, more preferably at least 1000, still more preferably at least 10,000, and most preferably at least 100,000 different compounds, or even more.

In the context of this invention, the term "small molecule" refers to compounds having molecular mass of less than 2000 Daltons, preferably less than 1500, still more preferably less than 1000, and most preferably less than 600 Daltons. Preferably but not necessarily, a small molecule is not an oligopeptide.

In a related aspect or in preferred embodiments, the invention provides a method of screening for potential antibacterial agents by determining whether any of a plurality of compounds, preferably a plurality of small molecules, is active on at least one target of a bacteriophage inhibitor protein or RNA. Preferred embodiments include those described for the above aspect, including embodiments which involve determining whether one or more test compounds bind to or reduce the level of activity of a bacterial target, and embodiments which utilize a plurality of different targets as described above.

The identification of bacteria-inhibiting phage ORFs and their encoded products also provides a method for identifying an active portion of such an encoded product. This also provides a method for identifying a potential antibacterial agent by identifying such an active portion of a phage ORF or ORF product. In preferred embodiments, the identification of an active portion involves one or more of mutational analysis, deletion analysis, or analysis of fragments of such products. The method can also include determination of a 3-dimensional structure of an active portion, such as by analysis of crystal diffraction patterns. In further embodiments, the method involves constructing or synthesizing a peptidomimetic compound, where the structure of the peptidomimetic compound corresponds to the structure of the active portion. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion that the peptidomimetic will interact with the same molecule as the phage protein and preferably will elicit at least one cellular response in common which relates to the inhibition of the cell by the phage protein.

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In preferred embodiments, the ORF or ORF product is or is derived or obtained from S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014 or product thereof.

The methods for identifying or screening for compounds or agents active on a bacterial target of a phage-encoded inhibitor can also involve identification of a phage-specific site of action on the target.

Preferably in the methods for identifying or screening for compounds active on such a bacterial target, the target is uncharacterized; the target is from an uncharacterized bacterium from Table 1; the site of action is a phage-specific site of action.

Further embodiments include the identification of inhibitor phage ORFs and bacterial targets as in aspects above.

An "active portion" as used herein denotes an epitope, a catalytic or regulatory domain, or a fragment of a bacteriophage inhibitor protein that is responsible for, or a significant factor in, bacterial target inhibition. The active portion preferably may be removed from its contiguous sequences and, in isolation, still effect inhibition.

By "mimetic" is meant a compound structurally and functionally related to a reference compound that can be natural, synthetic, or chimeric. In terms of the present invention, a "peptidomimetic," for example, is a compound that mimics the activity-related aspects of the 3-dimensional structure of a peptide or polyeptide in a non-peptide compound, for example mimics the structure of a peptide or active portion of a phage- or bacterial ORF-encoded polypeptide.

A related aspect provides a method for inhibiting a bacterial cell by contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein or RNA, where the target was uncharacterized. In preferred embodiments, the compound is such a protein, or a fragment or derivative thereof; a structural mimetic, e.g., a peptidomimetic, of such a protein or fragment; a small molecule; the contacting is performed in vitro, the contacting is performed in vivo in an infected or at risk organism, e.g., an animal such as a mammal or bird, for example, a human, or other mammal described herein; the bacterium is selected from a genus and/or species listed in Table 1; the bacteriophage inhibitor protein is uncharacterized; the bacteriophage inhibitor protein is from an uncharacterized phage listed in Table 1; the phage inhibitor protein is from one of S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

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In the context of targets in this invention, the term "uncharacterized" means that the target was not recognized as an appropriate target for an antibacterial agent prior to the filing of the present application or alternatively prior to the present invention. Such lack of recognition can include, for example, situations where the target and/or a nucleotide sequence encoding the target were unknown, situations where the target was known, but where it had not been identified as an appropriate target or as an essential cellular component, and situations where the target was known as essential but had not been recognized as an appropriate target due to a belief that the target would be inaccessible or otherwise that contacting the cell with a compound active on the target in vitro would be ineffective in cellular inhibition, or ineffective in treatment of an infection. Methods described herein utilizing bacterial targets, e.g., for inhibiting bacteria or treating bacterial infections, can also utilize "uncharacterized target sites", meaning that the target has been previously recognized as an appropriate target for an antibacterial agent, but where an agent or inhibitor of the invention is used which acts at a different site than that at which the previously utilized antibacterial agent, i.e., a phage-specific site. Preferably the phage-specific site has different functional characteristics from the previously utilized site. In the context of targets or target sites, the term "phage-specific" indicates that the target or site is utilized by at least one bacteriophage as an inhibitory target and is different from previously identified targets or target sites.

In the context of this invention, the term "bacteriophage inhibitor protein" refers to a protein encoded by a bacteriophage nucleic acid sequence which inhibits bacterial function in a host bacterium. Thus, it is a bacteria-inhibiting phage product.

In the context of this invention, the phrase "contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein" or equivalent phrases refer to contacting with an isolated, purified, or enriched compound or a composition including such a compound, but specifically does not rely on contacting the bacterial cell with an intact phage which encodes the compound. Preferably no intact phage are involved in the contacting.

Related aspects provide methods for prophylactic or therapeutic treatment of a bacterial infection by administering to an infected, challenged or at risk organism a therapeutically or prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein or RNA, or as described for the previous aspect.

Preferably the bacterium involved in the infection or risk of infection produces the identified target of the bacteriophage inhibitor protein or alternatively produces a homologous target compound. In preferred embodiments, the host organism is a plant or animal, preferably a mammal or bird, and more preferably, a human or other

mammal described herein. Preferred embodiments include, without limitation, those as described for the preceding aspect.

Compounds useful for the methods of inhibiting, methods of treating, and pharmaceutical compositions can include novel compounds, but can also include compounds which had previously been identified for a purpose other than inhibition of bacteria. Such compounds can be utilized as described and can be included in pharmaceutical compositions.

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In preferred embodiments of this and other aspects of the invention utilizing bacterial target sequences of a bacteriiophage inhibitory ORF product, the target sequence is encoded by a Staphylococcus nucleic acid coding sequence, preferably S. aureus, a Streptococcus nucleic acid coding sequence, preferably Streptococcus pneumoniae, or Enterococcus nucleic acid coding sequence. Possible target sequences are described herein by reference to sequence source sites.

The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. For the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage host genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

In the context of nucleic acid or amino acid sequences of this invention, the term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the homolog provides functionally equivalent biological function.

By "treatment" or "treating" is meant administering a compound or pharmaceutical composition for prophylactic and/or therapeutic purposes. The term "prophylactic treatment" refers to treating a patient or animal that is not yet infected but is susceptible to or otherwise at risk of a bacterial infection. The term "therapeutic treatment" refers to administering treatment to a patient already suffering from. infection.

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The term "bacterial infection" refers to the invasion of the host organism, animal or plant, by pathogenic bacteria. This includes the excessive growth of bacteria which are normally present in or on the body of the organism, but more generally, a bacterial infection can be any situation in which the presence of a bacterial population(s) is damaging to a host organism. Thus, for example, an organism suffers from a bacterial population when excessive numbers of a bacterial population are present in or on the organism's body, or when the effects of the presence of a bacterial population(s) is damaging to the cells, tissue, or organs of the organism.

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The terms "administer", "administering", and "administration" refer to a method of giving a dosage of a compound or composition, e.g., an antibacterial pharmaceutical composition, to an organism. Where the organism is a mammal, the method is, e.g., topical, oral, intravenous, transdermal, intraperitoneal, intramuscular, or intrathecal. The preferred method of administration can vary depending on various factors, e.g., the components of the pharmaceutical composition, the site of the potential or actual bacterial infection, the bacterium involved, and the infection severity.

The term "mammal" has its usual biological meaning referring to any organism of the Class Mammalia of higher vertebrates that nourish their young with milk secreted by mammary glands, e.g., mouse, rat, and, in particular, human, bovine, sheep, swine, dog, and cat.

In the context of treating a bacterial infection a "therapeutically effective amount" or "pharmaceutically effective amount" indicates an amount of an antibacterial agent, e.g., as disclosed for this invention, which has a therapeutic effect. This generally refers to the inhibition, to some extent, of the normal cellular functioning of bacterial cells that renders or contributes to bacterial infection.

The dose of antibacterial agent that is useful as a treatment is a "therapeutically effective amount." Thus, as used herein, a therapeutically effective amount means an amount of an antibacterial agent that produces the desired therapeutic effect as judged by clinical trial results and/or animal models. This amount can be routinely determined by one skilled in the art and will vary depending on several factors, such as the particular bacterial strain involved and the particular antibacterial agent used.

In connection with claims to methods of inhibiting bacteria and therapeutic or prophylactic treatments, "a compound active on a target of a bacteriophage inhibitor protein" or terms of equivalent meaning differ from administration of or contact with an intact phage naturally encoding the full-length inhibitor compound. While an intact phage may conceivably be incorporated in the present methods, the method at

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least includes the use of an active compound as specified different from a full length inhibitor protein naturally encoded by a bacteriophage and/or a delivery or contacting method different from administration of or contact with an intact phage encoding the full-length protein. Similarly, pharmaceutical compositions described herein at least include an active compound different from a full-length inhibitor protein naturally encoded by a bacteriophage or such a full-length protein is provided in the composition in a form different from being encoded by an intact phage. Preferably the methods and compositions do not include an intact phage.

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In accord with the above aspects, the invention also provides antibacterial 10 agents and compounds active on bacterial targets of bacteriophage inhibitor proteins or RNAs, where the target was uncharacterized as indicated above. As previously indicated, such active compounds include both novel compounds and compounds which had previously been identified for a purpose other than inhibition of bacteria. Such previously identified biologically active compounds can be used in embodiments of the above methods of inhibiting and treating. In preferred 15 embodiments, the targets, bacteriophage, and active compound are as described herein for methods of inhibiting and methods of treating. Preferably the agent or compound is formulated in a pharmaceutical composition which includes a pharmaceutically acceptable carrier, excipient, or diluent. In addition, the invention provides agents, 20 compounds, and pharmaceutical compositions where an active compound is active on an uncharacterized phage-specific site.

In preferred embodiments, the target is as described for embodiments of aspects above.

Likewise, the invention provides a method of making an antibacterial agent. 25 The method involves identifying a target of a bacteriophage inhibitor polypeptide or protein or RNA, screening a plurality of compounds to identify a compound active on the target, and synthesizing the compound in an amount sufficient to provide a therapeutic effect when administered to an organism infected by a bacterium naturally producing the target. In preferred embodiments, the identification of the target and 30 identification of active compounds include steps or methods and/or components as described above (or otherwise herein) for such identification. Likewise, the active compound can be as described above, including fragments and derivatives of phage inhibitor proteins, peptidomimetics, and small molecules. As recognized by those skilled in the art, peptides can be synthesized by expression systems and purified, or can be synthesized artificially. In preferred embodiments the inhibitory phage ORF 35 products is from S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus

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pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

As indicated above, sequence analysis of nucleotide and/or amino acid sequences can beneficially utilize computer analysis. Thus, in additional aspects the invention provides computer-related hardware and media and methods utilizing and incorporating sequence data from uncharacterized phage, e.g., uncharacterized phage listed in Table 1, preferably at least one of Staphylococcus aureus phage S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014, or 44 AHJD, Enterococcus sp. phage 182, or Streptococcus pneumoniae phage Dp-1. In general, such aspects can facilitate the above-described aspects. Various embodiments involve the analysis of genetic sequence and encoded products, as applied to the evaluating bacteriophage inhibitor ORFs and compounds and fragments related thereto. The various sequence analyses. as well as function analyses, can be used separately or in combination, as well as in preceding aspects and embodiments. Use in combination is often advantageous as the additional information allows more efficient prioritizing of phage ORFs for identification of those ORFs that provide bacteria-inhibiting function.

In one aspect, the invention provides a computer-readable device which includes at least one recorded amino acid or nucleotide sequence corresponding to one of the specified phage and a sequence analysis program for analyzing a nucleotide and/or amino acid sequence. The device is arranged such that the sequence information can be retrieved and analyzed using the analysis program. The analysis can identify, for example, homologous sequences or the indicated %s of the phage genome and structural motifs. Preferably the sequence includes at least 1 phage ORF or encoded product, more preferably at least 10%, 20%, 30%, 40%, 50%, 70%, 90%, or 100% of the genomic phage ORFs and/or equivalent cDNA, RNA, or amino acid sequences. Preferably the sequence or sequences in the device are recorded in a medium such as a floppy disk, a computer hard drive, an optical disk, computer random access memory (RAM), or magnetic tape. The program may also be recorded in such medium. The sequences can also include sequences from a plurality of different phage.

In this context, the term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the homolog provides functionally equivalent biological function.

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Similarly, the invention provides a computer analysis system for identifying biologically important portions of a bacteriophage genome. The system includes a data storage medium, e.g., as identified above, which has recorded thereon a nucleotide sequence corresponding to at least a portion of at least one uncharacterized bacteriophage genome, a set of program instructions to allow searching of the sequence or sequences to analyze the sequence, and an output device where the portion includes at least the sequence length as specified in the preceding aspect. The output device is preferably a printer, a video display, or a recording medium. More one than one output device may be included. For each of the present computer-related asepcts, the bacteriophage are preferably selected from the uncharacterized phage listed in Table 1, more preferably from bacteriophage 77, 3A, 96, 44 AHJD (S. aureus), Dp-1 (Streptococcus pneumoniae), or 182 (Enterococcus).

In keeping with the computer device aspects, the invention also provides a method for identifying or characterizing a bacteriophage ORF by providing a computer-based system for analyzing nucleotide or amino acid sequences, e.g., as describe above. The system includes a data storage medium which has recorded a sequences or sequences as described for the above devices, a set of instructions as in the preceding aspect, and an output device as in the preceding aspect. The method further involves analyzing at least one sequence, and outputting the analysis results to at least one output device.

In preferred embodiments, the analysis identifies a sequence similarity or homology with a sequence or sequences selected from bacterial ORFs encoding products with related biological function; ORFs encoding known inhibitors; and essential bacterial ORFs. Preferably the analysis identifies a probable biological function based on identification of structural elements or characteristic or signature motifs of an encoded product or on sequence similarity or homology. Preferably the uncharacterized bacteriophage is from Table 1, more preferably at least one of bacteriophage 77, 3A, 96, 44 AHJD (S. aureus), Dp-1 (Streptococcus pneumoniae), or 182 (Enterococcus). In preferred embodiments, the method also involves determining at least a portion of the nucleotide sequence of at least one uncharacterized bacteriophage as indicated, and recording that sequence on data storage medium of the computer-based system. In preferred embodiments, the analysis identifies a sequence similarity of homology with a S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

As used in the claims to describe the various inventive aspects and embodiments, "comprising" means including, but not limited to, whatever follows the word "comprising". Thus, use of the term "comprising" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present. By "consisting of" is meant including, and limited to, whatever follows the phrase "consisting of". Thus, the phrase "consisting of" indicates that the listed elements are required or mandatory, and that no other elements may be present. By "consisting essentially of" is meant including any elements listed after the phrase, and limited to other elements that do not interfere with or contribute to the activity or action specified in the disclosure for the listed elements. Thus, the phrase "consisting essentially of" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present depending upon whether or not they affect the activity or action of the listed elements.

Further embodiments will be apparent from the following Detailed Description and from the claims.

# BRIEF DESCRIPTION OF THE DRAWINGS

FIGURE 1A and 1B are flow schematics showing the manipulations used to convert pT0021, an arsenite inducible vector containing the luciferase gene, into pTHA or pTM, two *ars* inducible vectors. Vector pTHA contains BamH I, Sal I, and Hind III cloning sites and a downstream HA epitope tag. Vector pTM contains Bam HI and Hind III cloning sites and no HA epitope tag.

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FIGURE 2 is a schematic representation of the cloning steps involved to place the DNA segments of any of ORFs 17/ 19/ 43/ 102/104/182 or other sequences into pTHA to assess inhibitory potential. For subcloning into pTM or pT0021, Individual ORFs were amplified by the PCR using oligonucleotides targeting the ATG and stop codons of the ORFs. Using this strategy, Bam HI and Hind III sites were positioned immediately upstream or downstream, respectively of the start and stop codons of each ORF. Following digestion with Bam HI and Hind III, the PCR fragments were subcloned into the same sites of pT0021 or pTM. Clones were verified by PCR and direct sequencing.

FIGURE 3 shows a schematic representation of the functional assays used to characterize the bactericidal and bacteriostatic potential of all predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Fig. 3A) Functional assay on semi-solid support media. Fig. 3B) Functional assay in liquid culture.

FIGURE 4A, B, and C is a bar graph showing the results of a screen in liquid media to assess bacteriostatic or bactericidal activity of 93 predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Growth inhibition assays were performed as detailed in the Detailed Description. The relative growth of Staphylococcus aureus transformants harboring a given bacteriophage 77 ORF (identified on the bottom of the graph), in the absence or presence of arsenite, is plotted relative to growth of a Staphylococcus aureus transformant containing ORF 5, a non-toxic bacteriophage 77 ORF (which is set at 100%). Each bar represents the average obtained from three Staph A transformants grown in duplicate. Bacteriophage 77 ORFs showing significant growth inhibition consist of ORFs 17, 19, 102, 104, and 182.

FIGURE 5 shows a block diagram of major components of a general purpose computer.

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FIGURE 6 shows an ORF map for *Streptococcus pneumoniae* bacteriophage Dp-1 showing the ORF identifiers, genomic locations, and orientations of the 85 identified ORFs that were found to have ribosomal binding sites and thus are expected to be expressed.

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FIGURE 7 shows a schematic representation of the arsenite-inducible expression system present in a shuttle vector designed to express individual *Streptococcus* bacteriophage Dp-1 ORFs in *Streptococcus*. Various modifications can be readily made to such a vector, or other vectors can be readily constructed to provide inducible expression of ORFs in a particular host bacterium using well-known techniques.

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# DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention may be more clearly understood from the following description.

The tables will first be briefly described.

Table 1 is a listing of a large number of available bacteriophage that can be readily obtained and used in the present invention.

Table 2 shows the complete nucleotide sequence of the genome of Staphylococcus aureus bacteriophage 77.

Table 3 shows a list of all the ORFs from Bacteriophage 77 that were screened in the functional assay to identify those with anti-microbial activity.

Table 4 shows the predicted nucleotide sequence, predicted amino acid sequence, and physiochemical parameters of ORF 17/ 19/ 43/ 102/ 104/ 182]. These include the primary amino acid sequence of the predicted protein, the average molecular weight, amino acid composition, theoretical pI, hydrophobicity map, and predicted secondary structure map.

Table 5 shows homology search results. BLAST analysis was performed with ORFs 17/ 19/ 43/ 102/ 104/ 182 against NCBI non-redundant nucleotide and Swissprot databases. The results of this search indicate that: I) ORF 17 has no significant homology to any gene in the NCBI non-NCBI non-redundant nucleotide database, II) ORF 19 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 59 of bacteriophage phi PVL, III) ORF 43 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL, IV) ORF 102 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 38 of phi PVL, V) ORF 104 has no significant homology to any gene in the NCBI non-redundant nucleotide database, VI) ORF 182 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL.

Table 6 is a table from Alberts et al., MOLECULAR BIOLOGY OF THE CELL 3<sup>rd</sup> ed., showing the redundancy of the "universal" genetic code.

Table 7 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 3A.

Table 8 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 3A.

Table 9 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 96.

Table 10 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 96.

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Table 11 is a listing of sequences deposited in the NCBI public database (GeneBank) for bacteriophage listed in Table 1.

Table 12 is a listing of phage which encode a known lysis function, including the identified lysis gene.

Table 13 is a listing of bacteriophage which encode holin genes, where holin genes encode proteins which form pores and eventually enable other enzymes to kill the host bacterium.

Table 14 is a listing of bacteriophage which encode kil genes.

Table 15 is a list of *Staphylococcus aureus* sequences identified by accession number which may include sequences from genes coding for target sequences for the phage 77-encoded antimicrobial proteins or peptides. The sequences were obtained by searching GenBank for listings.

Table 16 shows the nucleotide sequence of the genome of *Staphylococcus* aureus phage 44 AHJD.

Table 17 lists and shows the sequence position of the 73 ORFs predicted to be encoded by *Staphylococcus aureus* bacteriophage 44 AHJD that are greater than 33 amino acids.

Table 18 shows the ORF sequences and putative amino acid sequences for the Staphylococcus aureus bacteriophage 44AHJD ORFs greater than 33 amino acids.

Table 19 shows the similarities in sequence identified between predicted Staphylococcus aureus bacteriophage 44 AHJD ORFs and sequences present in public databases.

Table 20 shows the homology alignments between predicted *Staphylococcus* aureus bacteriophage 44AHJD ORFs and the corresponding protein sequences present in public sequence databases.

Table 21 shows the complete nucleotide sequence of the genome of *Enterococcus* bacteriophage 182.

Table 22 lists and shows the sequence position of the 80 ORFs identified in bacteriophage 182 and that are greater than 33 amino acids.

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Table 23 shows the nucleotide and predicted amino acid sequence of all 80 ORFs identified in bacteriophage 182.

Table 24 shows the similarities identified to date in sequence between Enterococcus phage 182 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 25 shows the predicted amino acid sequence as well as the predicted secondary structures map for two *Enterococcus* bacteriophage 182 ORFs.

Table 26 shows the homology alignments between predicted *Enterococcus* bacteriophage 182 ORFs and the corresponding protein sequences present in public sequence databases.

Table 27 list *Enterococcus* sequences listed in GenBank providing possible Enterococcal target sequences for inhibitory *Enterococcus* bacteriophage 182 ORFs and other compounds with antibacterial activity.

Table 28 shows the complete nucleotide sequence of the genome of *Streptococcus* bacteriophage Dp-1.

Table 29 lists and shows sequence position of the 273 ORFs identified in Pneumococcal bacteriophage Dp-1 that are greater than 33 amino acids, 85 of which are predicted to be expressed in Dp-1 as having a ribosomal binding site. That set of 85 ORFs is shown in the attached drawings.

Table 30 shows the nucleotide and predicted amino acid sequence of all 273 ORFs identified in bacteriophage Dp-1 that are identified as being expressed.

Table 31 shows the similarities identified in sequence between *Streptococcus* phage Dp-1 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 32 shows the 4731 bp sequence of Dp-1 published by Sheehan et al., 1997).

Table 33 lists Streptococcus pneumoniae sequences listed in GenBank providing possible target sequences for inhibitory Streptococcus pneumoniae bacteriophage Dp-1 ORFs and other compounds with antibacterial activity

# Background:

As indicated above, the present invention is concerned, in part, with the use of bacteriophage coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents. Thus, the invention concerns the selection of relevant bacteria. Particularly relevant bacteria are those which are pathogens of a complex organism such as an animal, e.g., mammals,

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reptiles, and birds, and plants. Examples include Stapylococcus aureus, Enterococcus species, and Streptococcus pneumoniae. However, the invention can be applied to any bacterium (whether pathogenic or not) for which bacteriophage are available or which are found to have cellular components closely homologous to components targeted by phage of another bacterium.

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Thus, the invention also concerns the bacteriophage which can infect a selected bacterium. Identification of ORFs or products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such targets are thus identified as potential targets for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, a phage-encoded inhibitor can also inhibit such a homologous bacterial cellular component.

The demonstration that bacteriophage have adapted to inhibiting a host bacterium by acting on a particular cellular component or target provides a strong indication that that component is an appropriate target for developing and using antibacterial agents, e.g., in therapeutic treatments. Thus, the present invention provides additional guidance over mere identification of bacterial essential genes, as the present invention also provides an indication of accessability of the target to an inhibitor, and an indication that the target is sufficiently stable over time (e.g., not subject to high rates of mutation) as phage acting on that target were able to develop and persist. Thus, the present invention identifies a subset of essential cellular components which are particularly likely to be appropriate targets for development of antibacterial agents.

The invention also, therefore, concerns the development or identification of inhibitors of bacteria, in addition to the phage-encoded inhibitory proteins (or RNA transcripts), which are active on the targets of bacteriophage-encoded inhibitors. As described herein, such inhibitors can be of a variety of different types, but are preferably small molecules.

The following description provides preferred methods for use in the various aspects of the invention. However, as those skilled in the art will readily recognize, other approaches can be used to obtain and process relevant information. Thus the invention is not limited to the specifically described methods. In addition, the following description provides a set of steps in a particular order. That series of steps

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describes the overall development involved in the present invention. However, it is clear that individual steps or portions of steps may be usefully practiced separately, and, further, that certain steps may be performed in a different order or even bypassed if appropriate information is already available or is provided by other sources or methods.

## Selecting and Growing Phage, and Isolating DNA

Conceptually, the first step involves selecting bacterial hosts of interest. Preferably, but not necessarily, such hosts will be pathogens of clinical importance. Alternatively, because bacteria all share certain fundamental metabolic and structural features, these features can be targeted for study in one strain, for example a nonpathogenic one, and extrapolated to similarly succeed in pathogenic ones. Nonpathogenic strains may also exhibit initial advantages in being not only less dangerous, but also, for example, in having better growth and culturing characteristics and/or better developed molecular biology techniques and reagents. Consequently, advantageously the invention provides the ability target virtually any bacteria, but preferably pathogenic bacteria, with antimicrobial compounds designed and/or developed using bacteriophage inhibitory proteins and peptides from phage with non-pathogenic and/or pathogenic hosts.

We have selected Staphylococcus aureus, Streptococcus pneumoniae, various Enterococci, and Pseudomonas aeruginosa as initial exemplary pathogens. These bacteria are a major cause of morbidity and mortality in hospital-based infections, and the appearance of antibiotics resistance in all three organisms makes it increasingly difficult to treat benign infections involving these organisms. Such infections can include, for example, otitis media, sinusitis, and skin, and airway infections (Neu, H.C. (1992). Science 257, 1064-1073). However, the approach described below is clearly applicable to any human bacterial pathogens including but not restricted to Mycobacterium tuberculosis, Nesseria gonorrhoeae, Haemophilus influenza, Acinobacter, Escherichia coli, Shigella dysenteria, Streptococcus pyogenes, Helicobacter pylori, and Mycoplasma species. This invention can also be applied to the discovery of anti-bacterial compounds directed against pathogens of animals other than humans, for example, sheep, cattle, swine, dogs, cats, birds, and reptiles. Similarly, the invention is not limited to animals, but also applies to plants and plant pathogens.

In general, the bacteria are grown according to standard methodologies - employed in the art, including solid, semi-solid or liquid culturing, which procedures can be found in or extrapolated from standard sources such as Maloy, S.R., Stewart,

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V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press, or Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; or Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. Culture conditions are selected which are adapted to the particular bacterium generally using culture conditions known in the art as appropriate, or adaptations of those conditions.

Nucleic acids within these bacteria can be routinely extracted through common procedures such as described in the above-referenced manuals and as generally known to those skilled in the art. Those nucleic acid stocks can then be used to practice the other inventive aspects described below.

## Selection and Growth of Bacteriophage, and Isolation of DNA

The second step involves assembling a group of bacteriophages (phage collection) for one or more of the targeted bacterial hosts. While the invention can be utilized with a single bacteriophage for a pathogen or other bacterium, it is preferable to utilize a plurality of phage for each bacterium, as comparisons between a plurality of such phage provides useful additional information. Non-limiting examples of phage and sources for some of the above-mentioned pathogenic bacteria are found in Table 1. The criteria used to select such phages is that they are infectious for the microbe targeted, and replicate in, lyse, or otherwise inhibit growth of the bacterium in a measurable fashion. These phages can be very different from one another (representing different families), as judged by criteria such as morphology (head, tail, plate, etc.), and similarity of genome nucleotide sequence (cross-hybridization). Since such diverse bacteriophages are expected to block bacterial host metabolism and ultimately inhibit by a variety of mechanisms, their combined study will lead to the identification of different mechanisms by which the phages independently inhibit bacterial targets. Examples include degradation of host DNA (Parson K.A., and Snustad, D.P. (1975). J. Virol. 15, 221-444) and inhibition of host RNA transcription (Severinova, E., Severinov, K. and Darst, S.A. (1998). J.Mol. Biol. 279, 9-18). This, in turn, yields novel information on phage proteins that can inhibit the targeted microbe. As explained below, this 1) forms the basis of novel drug discovery efforts based on knowledge of the primary amino acid sequence of the phage inhibitor protein (e.g., peptide fragments or peptidomimetics) and/or 2) leads to the identification of bacterial biochemical pathways, the proteins of which are essential or significant for survival of the targeted microbe, and which enzymatic steps or

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chemical reactions can be targeted by classical drug discovery methods using molecular inhibitors, for example, small molecule inhibitors.

Bacteriophage are generally either of two types, lytic or filamentous, meaning they either outright destroy their host and seek out new hosts after replication, or else continuously propogate and extrude progeny phage from the same host without destroying it. Regardless of the phage life cycle and type, preferred embodiments incorporate phage which impede cell growth in measurable fashion and preferably stop cell growth. To this end, lytic phage are preferred, although certain nonlytic species may also suffice, e.g., if sufficiently bacteriostatic.

Various procedures that are commonly understood by those of skill in the art can be routinely employed to grow, isolate, and purify phage. Such procedures are exemplified by those found in such common laboratory aids such as Maloy, S.R., Stewart, V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press; Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; and Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. The techniques generally involve the culturing of infected bacterial cells that are lysed naturally and/or chemically assisted, for example, by the use of an organic solvent such as chloroform that destroys the host cells thereby liberating the phage within. Following this, the cellular debris is centrifuged away from the supernatant containing the phage particles, and the phage then subsequently and selectively precipitated out of the supernatant using various methods usually employing the use of alcohols and/or other chemical compounds such as polyethylene glycol (PEG). The resulting phage can be further purified using various density gradient/centrifugation methodologies. The resulting phage are then chemically lysed, thereby releasing their nucleic acids that can be conveniently precipitated out of the supernatant to yield a viral nucleic acid supply of the phage of interest.

Exemplary bacteriophage are indicated in Table 1, along with sources where those phage may be obtained.

Exemplary bacteria include the reference bacteria for the identified bacteriophage, available from the same sources.

## Characterizing Bacteriophage Genomes for ORFs

The third step involves systematically characterizing the genetic information contained in the phage genome. Within this genetic information is the sequence of all RNAs and proteins encoded by the phage, including those that are essential or

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instrumental in inhibiting their host. This characterization is preferably done in a systematic fashion. For example, this can be done by first isolating high molecular weight genomic DNA from the phage using standard bacterial lysis methods, followed by phage purification using density gradient ultracentrifugation, and extraction of nucleic acid from the purified phage preparation. The high molecular weight DNA is then analyzed to determine its size and to evaluate a proper strategy for its sequencing. The DNA is broken down into smaller size fragments by sonication or partial digestion with frequently cutting restriction enzymes such as Sau3A to yield predominantly 1 to 2 kilobase length DNA, which DNA can then be resolved by gel electrophoresis followed by extraction from the gel.

The ends of the fragments are enzymatically treated to render them suitable for cloning and the pools of fragments are cloned in a bacterial plasmid to generate a library of the phage genome. Several hundred of these random DNA fragments contained in the plasmid vector are isolated as clones after introduction into an appropriate bacterium, usually *Escherichia coli*. They are then individually expanded in culture and the DNA from each individual clone is purified. The nucleotide sequences of the inserts of these clones are determined by standard automated or manual methods, using oligonucleotide primers located on either side of the cloning site to direct polymerase mediated sequencing (e.g., the Sanger sequencing method or a modification of that method). Other sequencing methods can also be used.

The sequence of individual clones is then deposited in a computer, and specific software programs (for example, Sequencher<sup>TM</sup>, Gene Codes Corp.) are used to look for overlap between the various sequences, resulting in ordering of contig sequences and ultimately providing the complete sequence of the entire bacteriophage genome (one such example is given in Table 2 for *Staphylococcus aureus* bacteriophage 77; others are also provided herein). This complete nucleotide sequence is preferably determined with a redundancy of at least 3- to 5-fold (number of independent sequencing events covering the same region) in order to minimize sequencing errors.

Preferably, the bacterial strain used as a phage host should not possess any other innate plasmids, transposons, or other phage or incompatible sequences that would complicate or otherwise make the various manipulations and analyses more difficult.

Commercially available computer software programs are used to translate the nucleotide sequence of the phage to identify all protein sequences encoded by the — phage (hereafter called open reading frames or ORFs). (Customized software can clearly also be used.) As phages are known to transcribe their genome into RNA from

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both strands, in both directions, and sometimes in more than one frame for the same sequence, this exercise is done for both strands and in all six possible reading frames. As evolutionary constraints have forced the phage to conserve all of its vital protein sequences in as small a genome as possible, it is straightforward to identify all the proteins encoded by the phage by simple examination of the 6 translation frames of the genome. Once these ORFs are identified, they are cataloged into a phage proteome database (Table 3 lists ORFs identified from phage 77; ORF lists are also provided for other exemplary phage). This analysis is preferably performed for each phage under study. The process of ORF identification can be varied depending on the desired results. For example, the minimum length for the putative encoded polypeptide can be varied, and/or putative coding regions that have an associated Shine-Dalgarno sequence can be selected. In the case of phage 77 ORFs, such parameter adjustment was performed and resulted in the identification of ORFs as listed herein. Different parameters had resulted in the identification of the ORFs listed in the preceding U.S. Provisional Application 60/110,992, filed December 3, 1998, which is hereby incorporated by reference in its entirety.

Exemplary phage 77 ORFs identified in that provisional application and as identified herein are shown in the following table:

ORF ID from 60/110,992	Genomic position	a.a. size	Start codon	ORF ID from 241/190	Genomic position	a.a. size	Start codon
77ORF016	2369-24024	251	TTG	77ORF017	23269-23982	237	ATG
77ORF019	39845-40501	218	ATA	77ORF019	39851-40501	216	ATG
77ORF050	29268-29564	98	ATG	77ORF182	29268-29564	98	ATG
77ORF050	29268-29564	98	ATG	77ORF043	29304-29564	86	ATG
77ORF067	34312-34551	79	CTG	77ORF104	34393-34551	52	ATG
77ORF146	29051-29212	53	ATG	77ORF102	29051-29212	53	ATG

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# Identifying and Characterizing Inhibitory Phage ORFs

The fourth step entails identifying the phage protein or proteins or RNA transcripts that have the ability to inhibit their bacterial hosts. This can be accomplished, for example, by either or both of two non-mutually exclusive methods. The first method makes use of bioinformatics. Over the past few years, a large amount of nucleotide sequence information and corresponding translated products have become available through large genome sequencing projects for a variety of organisms including mammals, insects, plants, unicellular eukaryotes (yeast and fungi), as well as several bacterial genomes such as E. coli, Mycobacterium tuberculosis, Bacillus subtilis, Staphylococcus aureus and many others. Such sequences have been deposited in public databases (for example, non-redundant

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sequence database at GenBank and SwissProt protein sequence database) (http://www.ncbi.nlm.nih.gov)) and can be freely accessed to compare any specific query sequence to those present in such databases. For example, GenBank contains over 1.6 billion nucleotides corresponding to 2.3 million sequence records. Several computer programs and servers (e.g., TBLASTN) have been created to allow the rapid identification of homology between any given sequence from one organism to that of another present in such databases, and such programs are public and available free of charge.

In addition, it has been well established that basic biochemical pathways can be conserved in very distant organisms (for example bacteria and man), and that the proteins performing the various enzymatic steps in these pathways are themselves conserved at the amino acid sequence level. Thus, proteins performing similar functions (e.g. DNA repair, RNA transcription, RNA translation) have frequently preserved key structural signatures, identifiable by similarities across regions of proteins (domains and motifs). The antimicrobials of the present invention will preferably target features and targets that are highly characteristic or conserved in microbes, and not higher organisms.

Most genomes encode individual proteins or groups of proteins that can be assembled into protein families that have been evolutionarily conserved. Therefore, similarity between a new query sequence and that of a member of a protein family (reference sequences from public databases) can immediately suggest a biochemical function for the novel query sequence, which in our case is a phage ORF.

The sequence homology between individual members of evolutionarily distant members of a protein family is usually not randomly distributed along the entire length of the sequence but is often clustered into "motifs" and "domains". These correspond to key three-dimensional folds that form key catalytic and/or regulatory structures that perform key biochemical function(s) for the group of proteins. Commercially available computer software programs can identify such motifs in a new query sequence, again providing functional information for the query sequence. Such structural and functional motifs have also been derived from the combined analysis of primary sequence databases (protein sequences) and protein structure databases (X-ray crystallography, nuclear magnetic resonance) using so-called "threading" methods (Rost B,l and Sander C. (1996). Ann. Rev. Biophy. Biomol. Struct. 25, 113-136).

Such motifs and folds are themselves deposited in public databases which can be directly accessed (for example, SwissProt database; 3D-ALI at EMBL, Heidelberg; PROSITE). This basic exercise leads to a structural homology map in which each of

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the phage ORFs has been probed for such similarities, and where initial structural and functional hits are identified (selected examples of sequence homologies detected between individual ORFs from the genome of *Staphylococcus aureus* bacteriophage 77 and sequences deposited in public databases are shown in Table 5 for ORFs 17/19/43/102/104/182).

This analysis can point out phage proteins with similarity to proteins from other phages (such as those for *E. coli*) playing an important role in the basic biochemical pathways of the phage (such as DNA replication, RNA transcription, tRNAs, coat protein and assembly). Selected examples of such proteins include integrase and capsid protein. Therefore, this analysis enables identification and elimination of non-essential ORFs as candidates for an inhibitor function, as well as the identification of (potentially) useful ones.

In addition, this analysis can point out specific ORFs as possible inhibitor ORFs. For example these ORFs may encode proteins or enzymes that alter bacterial cell structure, metabolism or physiology, and ultimately viability. Examples of such proteins present in the genome of *Staphylococcus aureus* bacteriophage 77 include orf14 (deoxyuridine triphosphatase from bacteriophage T5), and orf15 (sialidase). (These ORF identifications are as listed in provisional application 60/110,992.) Other examples include ORFs 9 and 12 of *S. aureus* phage 44 AHJD, which encode the putative lysis functions found in many bacteriophages – a "holin" and an "amidase".

In addition, it is well known that bacterial and eukaryotic viruses can usurp pathways from their host in order to use them to their advantage in blocking host cellular pathways upon infection. The phage can achieve this by 1) directly producing an inhibitor of a key host pathway (e.g. T7 gene 0.5 and 2), 2) directly producing a novel activity (e.g. T4 DNA polymerase), and 3) altering concentrations of cell components by producing similar functions (e.g. T4 transfer RNAs). The identification of sequence similarity between phage ORFs and bacterial host genome sequences will be highly indicative of such a mechanism. (Selected examples of such homologies are listed in Figure 4 of the provisional application 60/110,992 and include orf4 (homologous to autolysin), orf20 (hypothetical protein from Staphyloccus aureus.)) These ORFs can be analyzed by a standard biochemical approach to directly test their inhibitor functions (e.g., as described below).

Alternatively, a homology search may reveal that a given phage ORF is related to a protein present in the databases having an activity known to be inhibitory, ( $e.\overline{g}$ . inhibitor of host RNA polymerase by  $E.\ coli$  bacteriophage T7. Such a finding would implicate the phage ORF product in a related activity. This will also suggest that a

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new antimicrobial could be derived by a mimetic approach (e.g., peptidomimetic) imitating this function or by a small molecule inhibitor to the bacterial target of the phage ORF, or any steps in the relevant host metabolic pathway, e.g., high throughput screening of small molecule libraries. Selected examples of such similarity between ORFs of Staphyloccus aureus bacteriophage 77 and proteins with inhibitor functions for bacterial hosts are listed in Figure 4 of the provisional application 60/110,992. These include orf9 (similar to bacteriophage P1 kilA function), and orf4 (autolysin of Staphylococcus aureus, amidase enzymatic activity).

A reason for the biochemical study of individual ORFs for inhibitor function is that their expression or overexpression will block cellular pathways of the host, ultimately leading to arrest and/or inhibition of host metabolism. In addition, such ORFs can alter host metabolism in different ways, including modification of pathogenicity. Therefore, individual ORFs identified above are expressed, preferably overexpressed, in the host and the effect of this expression or overexpression on host metabolism and viability is measured. This approach can be systematically applied to every ORF of the phage, if necessary, and does not rely on the absolute identification of candidate ORFs by bioinformatics. Individual ORFs are resynthesized from the phage genomic DNA, e.g., by the polymerase chain reaction (PCR), preferably using oligonucleotide primers flanking the ORF on either side. These single ORFs are preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial host such as E. coli, but containing the necessary information for plasmid replication in the target microbe such as S. aureus (hereafter referred to as shuttle vector). Shuttle vectors and their use are well known in the art.

Such shuttle vectors preferably also contain regulatory sequences that allow inducible expression of the introduced ORF. As the candidate ORF may encode an inhibitor function that will eliminate the host, it is beneficial that it not be expressed prior to testing for activity. Thus, screening for such sequences when expressed in a constitutive fashion is less likely to be successful when the inhibitor is lethal. In the exemplary inducible system presented in Figure 1A, 1B, 2, and 7, regulatory sequences from the ars operon of S. aureus are used to direct individual ORF expression in S. aureus (or other bacteria in which the ars system is functional). The ars operon encodes a series of proteins which normally mediate the extrusion of arsenite and other trivalent oxyanions from the cells when they are exposed to such toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are

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present. (Tauriainen, S. et al. (1997) App. Env. Microb., Vol. 63, No. 11, p. 4456-4461.)

Therefore, individual phage ORFs can be expressed in *S. aureus* in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *S. aureus* clones expressing such individual phage ORFs. Toxicity of the phage inhibitor ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reduced or arrested host metabolism can be measured by pulse-chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis. Similar constructs can be made and used for other bacteria using well-known techniques.

Those skilled in the art are familiar with a variety of other inducible systems which can also be used for the controlled expression of phage ORFs, including, for example, lactose (see *e.g.*, Stratagene's LacSwitch™II system; La Jolla, CA) and tetracycline-based systems (see, *e.g.* Clontech's Tet On/Tet Off™ system; Palo Alto, CA). The arsenite-inducible system described is further depicted in Figures 1, 2 and 7.

The selection or construction of shuttle vectors and the selection and use of inducible systems are well known and thus other shuttle vectors appropriate for other bacteria can be readily provided by those skilled in the art, e.g., for use in other bacterial species.

Standard methodologies for expressing proteins from constructs, and isolating and manipulating those proteins, for example in cross-linking and affinity chromatography studies, may be found in various commonly available and known laboratory manuals. See, e.g., Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J., and Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.

It has been found that certain phage or other viruses inhibit host cells, at least in part, by producing an antisense RNA which binds to and inhibits translation from a bacterial RNA sequence. Thus, in the case of potentially inhibitor RNA transcripts encoded by the phage genome, a strong indicator of a possible inhibitory function is provided by the identification of phage sequence which is the identical to or fully complementary (or with only a small percentage of mismatch, e.g., <10%, preferably less than 5%, most preferably less than 3%, to a bacterial sequence. This approach is convenient in the case of bacteria that have been essentially completely sequenced, as the comparison can be performed by computer using public database information.

The inhibitory effect of the transcript can be confirmed using expression of the phage sequence in a host bacterium. If needed, such inhibitory can also be tested by transfecting the cells with a vector that will transcribe the phage sequence to form RNA in such manner that the RNA produced will not be translated into a polypeptide. Inhibition under such conditions provides a strong indication that the inhibition is due to the transcript rather than to an encoded polypeptide.

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In an alternative, the expression of an ORF in a host bacterium is found to be inhibitory, but the inhibition is found to be due to an RNA product of the genomic coding region. For antisense inhibition, the sequence of the bacterial target nucleic acid sequence can be identified by inspection of the phage sequence, and the full sequence of the relevant coding region for the bacterial product can be found from a database of the bacterial genomic sequence or can be isolated by standard techniques (e.g., a clone in a genomic library can be isolated which contains the full bacterial ORF, and then sequenced).

In either case, the identification of a target which is inhibited by an RNA transcript produced by a phage provides both the possible inhibition of bacteria naturally containing the same target nucleic acid sequence, as well as the ability to use the target sequence in screening for other types of compounds which will act directly on the target nucleic acid sequence or on a polypeptide product expressed or regulated, at least in part, by the target of the inhibitory phage RNA.

In some cases it will be found that the target of an inhibitory phage RNA or protein has previously been found to be a target of an inhibitory phage RNA or protein has previously been found to be a target for an antibacterial agent. In such cases, the phage inhibitor can still provide useful information if it is found that the phage-encoded product acts at a different site than the previously identified antibacterial agent or inhibitor, i.e., acts at a phage-specific site. For many targets, action at a different site provides highly beneficial characteristics and/or information. For example, an alternate site of inhibitor action can at least partially overcome a resistance mechanism in a bacterium. As an illustration, in many cases, resistance is due, in large part, to altered binding characteristics of the immediate target to the antibacterial agent. The altered binding is due to a structural change which prevents or destabilizes the binding. However, the structural change is frequently quite local, so that compounds which bind at different local sites will b unaffected or affected to a much lesser degree. Indeed, in some cases the local sites will be on a different molecule and so may be completely unaffected by the local structural change creating resistance to the original agent(s). An example of resistance due to altered binding is

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provided by methicillin-resistant *Staphylococcus aureus*, in which the resistance is due to an altered penicillin-binding protein.

In other cases, a new site of action can have improved accessibility as compared to a site acted on by a previously identified agent. This can, for example, assist in allowing effective treatment at lower doses, or in allowing access by a larger range of types of compounds, potentially allowing identification of more potential active agents.

Another advantage is that the structural characteristics of a different site of action will lead to identification and/or development of inhibitors with different structures and different pharmacological parameter. This can allow a greater range of possibilities when selecting an antibacterial agent.

Yet further, different sites often produce different inhibitory characteristics in the target organism. This is commonly the case for multi-domain target proteins. Thus, inhibition targeting an alternate site can produce more efficacious action, e.g., faster killing, slower development of resistance, lower numbers of surviving cells, and different secondary effects (for example, different nutrient utilization).

#### Staphylococcus aureus phage 77

As indicated above, the present invention is concerned, in part, with the use of bacteriophage 77 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

As described, phage 77 ORFs 17, 19, 43, 102, 104, and 182 have been found to have bacteria inhibiting function. Identification of ORFs 17, 19, 43, 102, 104, and 182 and products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such a target is thus identified as a potential target for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, an inhibitor encoded by phage 77 ORF 17, 19, 43, 102, 104, or 182 can also inhibit such a homologous bacterial cellular component.

Possible bacterial target sequences are described herein by reference to sequence source sites. In preferred embodiments, the sequence encoding the target corresponds

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to a S. aureus nucleic acid sequence available from numerous sources including S. aureus sequences deposited in GenBank, S. aureus sequences found in European Patent Application No. 97100110.7 to Human Genome Sciences, Inc. filed January 7, 1997, S. aureus sequences available from TIGR at

http://www.tigr.org/tdb/mdb/mdb.html, and S. aureus sequences available from the Oklahoma University S. aureus sequencing project at the following URL: <a href="http://www.genome.ou.edu/staph\_new.html">http://www.genome.ou.edu/staph\_new.html</a>. Such possible targets are particularly applicable to S aureus phages 77, 3A, 96, and 44 AHJD.

The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a S. aureus coding sequence corresponding to a sequence listed in Table 15 herein. The listing in Table 15 describes S. aureus sequences currently listed with GenBank. Again, for the sake of brevity, the sequences are described by reference to the database accession numbers instead of being written out in full herein. In cases where an entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage host S. aureus genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

#### Staphyloccus aureus phage 44 AHJD

The present invention also can utilize the identification of naturally occurring DNA sequence elements within *Staphylococcus aureus* bacteriophage 44AHJD which encode proteins with antimicrobial activity.

Such identification can utilize bioinformatics identification of specific proteins (ORFs) utilized by Staphylococcus aureus bacteriophage 44AHJD during the viral life cycle, resulting in a slowing or arrest of growth of the bacterial host, or in death, of the Staphylococcus aureus host including lysis of the infected bacteria. Thus, some of the bacteriophage 44AHJD DNA sequences encoding these proteins (ORFs) are predicted to encode antimicrobial functions. Information derived from these DNA sequences and translated ORFs can, in turn, be utilized to develop inhibitory compounds by peptidomimetics that can also function as antimicrobials. In addition, the identification of the host bacterial proteins that are targeted and inhibited by the

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antimicrobial bacteriophage ORFs can themselves provide novel targets for drug discovery.

The methodology described above is used to identify and characterize DNA sequences from *Staphylococcus* sp. bacteriophage 44 AHJD that have antimicrobial activity. As described in the Examples, the *Staphylococcus aureus* propagating strain (PS 44A), obtained from the Felix d'Herelle Reference Centre (#HER 1101), was used as a host to propagate its phage 44AHJD, also obtained from the Felix d'Herelle Reference Centre (#HER 101). By sequencing, we found that bacteriophage 44AHJD consists of 16,668 bp (Table 16) predicted to encode 73 ORFs greater than 33 amino acids (Tables 17 & 18). Computational analysis of the predicted protein products of *Staphylococcus aureus* bacteriophage 44AHJD identified homolgs in public sequence databases as listed inTable 19 and 20, along with the accompanying list of related proteins.

From this analysis, it is apparent that 3 genes (ORF 3, 7, and 8) are related to structural proteins found in other bacteriophages. These include genes predicted to encode a tail protein (ORF 3), an upper collar/connector protein of the phage virion (ORF 7), and a lower collar protein (ORF 8). Bioinformatics has also identified one gene whose product is likely involved in phage DNA synthesis. One gene (ORF 1) shows significant homology to DNA polymerases of a number of bacteriophages, bacteria and fungi, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 44AHJD. ORF 2 encodes a protein with homology to the dinC gene of Bacillus subtilis that encodes a protein involved in teichoic acid biosynthesis. Teichoic acid is a polyphosphate polymer found in some, but not all, Gram positive organisms (and not in Gram negative organisms), where it is attached to the peptidoglycan layer. The phage protein may thus be involved in the synthesis of this material for incorporation into the cell wall, allowing enhanced lysis by the phage lysis enzymes or, as many enzymes can function in "reverse reactions", may be involved in its degradation allowing for penetration of the peptidoglycan and phage genome entry into the cell following adsorption. The similarity between Staphylococcus aureus bacteriophage 44AHJD and E. coli phage T7 indicates that they may share similar mechanisms of replication and growth. Both phages belong to the Pododviridae Family of bacteriophages and are members of the "T7-like" Genus of this Family (Ackermann and DuBow; VIth ICTV Report).

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Two genes, ORF 9 and 12, were identified with the potential to encode antimicrobial protein products. The homology alignments are shown in Tables 19 and 20. The predicted product of ORF 9 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms, including that from the Staphylococcus aureus bacteriophage Twort. ORF 12 of Staphylococcus aureus bacteriophage 44AHJD shows homology to a set of lysis proteins from several bacteriophages. These lysis proteins are also referred to as holins, and represent phage-encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the cell wall and thus lyse the bacterium.

Thus, in particular embodiments, the present invention provides a nucleic acid sequence isolated from Staphylococcus aureus bacteriophage 44AHJD comprising at least a portion of one of the genes described above with antimicrobial activity. For example, ORF 1 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORF 9 directly encodes a polypeptide with antimicrobial activity. ORF 9 is predicted to encode an amidase, a protein known to act as a cell wall degrading enzyme. ORF 12 likely encodes a holin function required for transit of the phage amidase (gene 9 product) to the periplasm. When this type of gene product from Bacillus phage phi 29 (gene 14), was cloned in Escherichia coli, cell death ensued (Steiner et al., 1993). Thus, production of proteins from Bacillus phage phi 29 gene 14 in E. coli resulted in cell death, whereas production of protein from Bacillus phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner et al., 1993).

The present invention also provides the use of the Staphylococcus

30 bacteriophage 44 AHJD antimicrobial ORFs or ORF products as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from Staphylococcus bacteriophage 44 AHJD killer ORFs.

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#### Enterococcus phage 182

Bacteriophage 182 was obtained from the Felix D'Herelle phage collection (Ste. Foy, Quebec) and infects *Enterococcus sp.* Group D. The genome of *Enterococcus* bacteriophage 182 consists of 17,833 bp (Table 21) and is predicted to encode 80 ORFs greater than 33 amino acids (Tables 22 and 23). Computational analysis of the predicted protein products of *Enterococcus* bacteriophage 182 was performed in order to identify protein products related to those deposited in public databases. Bacteriophage 182 protein products which detected sequences with significant sequence similarity in public databases are listed in Table 24 and 26, along with the accompanying list of related proteins.

From this analysis, it is apparent that 5 genes (ORF 001, 004, 007, 009, and 011) are related to structural proteins of several *Bacillus* phages – *Bacillus* bacteriophage PZA, phi-29, and B103. These include genes predicted to encode a tail protein (ORF 001), a head protein (ORF 004), and upper collar protein (ORF 007), a lower collar protein (ORF 009), and a pre-neck appendage protein (ORF 011). Two gene products are predicted to encode genes which direct phage morphogenesis – these are ORF 005 and 019.

Bioinformatics has also identified three genes whose products are likely involved in phage DNA synthesis. One gene, ORF 002 shows significant homology to DNA polymerases of a number of bacteriophages, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 182. ORF 006 encodes a protein with homology to the encapsidation proteins of several other bacteriophages, including *Bacillus* phage phi-29 (P11014), PZA (P07541), and B103 (X99260) and *Streptococcus* phage CP-1 (Z47794). These gene products catalyze the *in vivo* and *in vitro* genome-encapsidation reaction (Garvey et al., 1985). Proteins involved in genome packaging have been shown to have additional activities that affect biochemical reactions in other phages and their hosts. For example, the coat protein of the RNA bacteriophage MS2 interacts with viral RNA to translationally repress replicase synthesis (Pickett and Peabody, 1993). This protein-RNA interaction also plays a role in genome encapsidation, enveloping a single copy of the viral genome in a protein shell composed of many molecules of coat protein. In addition, the bacteriophage λ terminase enzyme can be lethal to *E. coli* when expressed,

suggesting cleavage of packaging sites in the bacterial chromosome. Also present within bacteriophage 182 is a gene, ORF 010, that encodes a protein that is related to the terminal proteins of *Bacillus* phage Nf (P06812), *Bacillus* phage GA-1 (X96987) and *Bacillus* phage B103 (X99260). DNA terminal proteins are linked to the 5' ends of both strands of the genome and are essential for DNA replication playing a role in initial priming of DNA replication. The similarity between *Enterococcus* bacteriophage 182 and Bacillus phages phi-29, PZA, and B103 indicates that they may share similar mechanisms of replication and growth. Protein-primed DNA replication is a well described phenomenon, and in the phi-29-like phages, the ends of the DNA serve as origins and termini of replication (Gutiérrez et al., 1986; Yoshikawa et al., 1985).

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There is also a gene (ORF 015) that encodes a protein showing homology to an early protein product of *Bacillus* bacteriophage PZA and the single-strand nucleic acid binding protein of bacteriophage B103.

Two genes, ORF 008 and 014, were identified with the potential to encode anti-microbial protein products. The homology alignments are shown in Tables 24 & 26 and biochemical features of the predicted polypeptides shown in Table 25. The predicted product of ORF 008 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms. ORF 014 of Enterococcus 182 shows homology to a set of lysis proteins from Bacillus bacteriophage phi-29, PZA, and B103. These lysis proteins are also referred to as holins and represent phage encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the outer cell wall and thus lyse the bacterium.

Thus, the present invention provides a nucleic acid sequence obtained from *Enterococcus* bacteriophage 182 comprising at least a portion of a phage 182 ORF, preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 002 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORFs 008 or 014 directly encode polypeptides with anti-microbial activity. ORF 008 is predicted to encode an

autolytic lysozyme, a protein known to have anti-microbial activity (Martin et al., 1998). ORF 014 likely encodes a holin function required for transit of the phage murein hydrolases to the periplasm. When the related product from Bacillus phage phi 29 (gene 14), was cloned in Escherichia coli, cell death ensued (Steiner et al., 1993). Thus, production of proteins from Bacillus phage phi 29 gene 14 in E. coli resulted in cell death, whereas production of protein from Bacillus phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner et al., 1993).

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The present invention also provides the use of the Enterococcus bacteriophage 182 anti-microbial ORFs as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from Enterococcus bacteriophage 182 killer ORFs. This can be done where the structure of the peptidomimetic compound corresponds to the structure of the active portion of a product of an ORF. In this analysis, the peptide backbone is transformed into a carbon based hydrophobic structure that can retain cytostatic or cytocidal activity for the bacterium. This is done by standard medicinal chemistry methods, measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These mimetics also represent lead compounds for the development of novel antibiotics. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion of a product of one of the Enterococcus ORFs listed, that the peptidomimetic will interact with the same molecule as the product of the ORF, and preferably will elicit at least one cellular response in common which relates to the inhibition of the cell by the phage protein.

To validate the identity of an ORF as a killer ORF, it is preferably expressed in the host or other test bacterial organism and the effect of this expression on bacterial growth and replication is assessed. Therefore, all individual ORFs identified herein, e.g., those identified above, can be expressed, preferably overexpressed, in a suitable host bacterium e.g., a host *Enterococcus* and the effect of this expression or overexpression on host metabolism and viability can be measured.

Individual ORFs can be resynthesized from the phage genomic DNA by the polymerase chain reaction (PCR) using oligonucleotide primers flanking the ORF on

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either side. Those skilled in the art are familiar with the design and synthesis of appropriate primer sequences. These single ORFs are preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial host such as *E. coli*, but containing the necessary information for plasmid replication in the target microbe, *Enterococcus* sp. (hereafter referred to as a shuttle vector).

This shuttle vector also preferably contains regulatory sequences that allow inducible expression of the introduced ORF. As the candidate ORF may encode a killer function that will eliminate the host, it is highly advantageous that it not be expressed (or at least not expressed at a substantial level) prior to testing for activity; thus screening for such sequences in a constitutive fashion is less likely to be successful (lethality). In an example presented in Fig. 7, regulatory sequences from the ars operon are used to direct individual ORF expression in Enterococcus. The ars operon encodes a series of proteins which normally mediate the extrusion of arsenite and several other trivalent oxyanions from the cells when they are exposed to such toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are present.

Therefore, individual phage ORFs can be expressed in *Enterococcus* or other suitable host in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *Enterococcus* (or other host cells) clones expressing such individual phage ORFs. Toxicity of the phage killer ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reducing or arresting host metabolism can be measured by pulse chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis.

Of course, other inducible regulatory sequences (e.g., promoters, operators, etc.) may be used (e.g., systems using positive induction of expression or systems using release of repression). A variety of such systems are known to those-skilled in the art and can be utilized in the present invention.

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Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art. In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Having the phage 182 ORFs, e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described, other anti-microbial sequences from other bacteriophage sources can be identified and isolated using methods described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

The invention also provides bacteriophage anti-microbial DNA segments from other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences which are highly homologous. The bacteriophage anti-microbial DNA segment from bacteriophage 182 can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on nucleic acid and/or amino acid sequence comparisons. As with the phage 182 inhibitory sequences, such homologous coding sequences and products can be used as antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

Enterococcus sequences are listed in Table 27 by accession number, providing identification of possible targets of Enterococcus phage inhibitory ORF products, e.g., from phage 182.

## Streptococcus pneumoniae

As indicated in the Summary above, the present invention is concerned with the use of *Streptococcus* sp. bacteriophage Dp-1 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

Streptococcus pneumoniae is an important cause of community-acquired pneumonia and a major cause of otitis media, sinusitis, and meningitis in children and adults. In Spain and other Mediterranean countries, the majority of S. pneumoniae are relatively resistant to penicillin (Klugman, 1990; Fenoll et al., 1991; Jorgensen et al., 1990). These strains also have decreased susceptibility to broad-spectrum cephaloporins, which are frequently used in the empiric treatment of meningitis and

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other serious invasive bacterial infections. High-level resistance of pneumococci has been encountered in Hungary where 70% of children who were colonized with *S. pneumoniae* carried penicillin resistant strains that were also resistant to tetracycline, erythromycin, trimethoprim/sulfamethoxazole, and 30% resistant to chloramphenicol (Neu, 1992). The resistance of pneumococci to macrolides such as erythromycin averages 20-25% in France, ~20% in Japan, and <10% in Spain (Neu, 1992).

The antimicrobial susceptibilities and distribution of serotypes of the 42 isolates of *S. pneumoniae* in southern Taiwan from invasive infections have been recently determined (Hseuh et al., 1996). Resistance rates among these isolates were: erythromycin, 61.9%; clindamycin, 47.6%; chloramphenicol, 19%; and tetracycline, 73.8%. Resistance to three or more classes of antibiotics was found in 33.3% of the isolates. Bacteremic pneumonia and primary bacteremia accounted for 64.3% of the infections and mortality was 42.6%. Given the severity of these infections despite adequate antibiotic therapy, there is clearly a need for introduction of new therapeutic options to prevent mortality due to invasive *S. pneumoniae* infections.

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Pneumococcal phages belong to four families and they present a great variety in morphology, including lytic and temperate phages (for a review, see Garcia et al., 1997). Examples of lytic phages are Cp-1 and Dp-1, whereas examples of temperate phages are HB-3, EJ-1, and HB-746. The complete nucleotide sequence and functional organization of Cp-1 has been reported (Martin et al., 1996). Cp-1 has a 19,345 bp double-stranded DNA genome, with a terminal protein covalently linked to its 5' ends, that replicates by a protein primed mechanism. The phage contains 29 ORFs, 23 on one strand and 6 on the opposite. When these predicted proteins were compared to sequences compiled in GenBank EMBL databases, to ORFs showed significant similarity to proteins of bacteriophage 29 that infects B. subtilis (Martin et al., 1996). The similar proteins corresponded to those involved in DNA replication (terminal protein and DNA polymerase), structural and morphogenic proteins (major head, collar, connector, tail, and encapsidation proteins), and proteins involved in lysis function (holin and lysozyme). In its strategy of lysis, the holin gene product inserts itself into the cell membrane, allowing access of the lysozyme to the peptidoglycan. Expression of the Cp-1 holin protein in E. coli results in cell death after 2-hours of induction, but did not lead to lysis (Garcia et al., 1997). Cells harboring a plasmid construction with holin and lysozyme genes together did lyse after induction and the

viability loss was similar to that of the culture expressing holin alone. Cloning of these lytic genes in *S. pneumoniae* showed that both genes had the same effect as in *E. coli*. That is, holin itself did not lyse the culture but the viability loss was noticeable, whereas both holin and lysozyme together were capable of lysing M31, an amidase deleted mutant (Garcia et al., 1997).

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Recently, a small portion (~4 kbp) of a second *S. pneumoniae* phage, Dp-1, has been sequenced (Sheehan et al., 1997). This portion contains the genes coding for the lytic system (Sheehan et al., 1997) and shows a modular organization similar to that described for Cp-1. However, in this case, a single chimeric protein appears to be made in which the N-terminal domain is highly similar to that of the murein hydrolase coded by a gene found in the phage BK5-T that infects *Lactococcus lactis*, and the C-terminal domain is homologous to holins. Thus, both functions appear to have been combined in a novel chimeric protein.

Bacteriophage Dp-1 was obtained from Dr. P. Garcia (Departamento de Microbiologia Molecular, Centro de Departamento de Investigaciones Biologicas, Consejo Superior de Investigaciones Cientificas, Velazquez, Madrid, Spain). We found that Dp-1 has a double-stranded DNA genome of 56,506 bp, predicted to encode 85 ORFs greater than 33 amino acids and with upstream Shine-Dalgarno motifs for translation initiation (Tables 28 & 30, and Fig. 6). Computational analysis of the predicted protein products of *Streptococcus* bacteriophage Dp-1 protein products, which detected homologs in public databases, are listed inTable 31, along with the accompanying list of related proteins.

From this analysis, it is apparent that several predicted genes of Dp-1 encode polypeptides that are related to structural proteins. ORFs 001, 002, 004, and 030 are predicted to encode tail proteins, minor structural proteins, and minor capsid proteins (Table 31). We also note the identification of several gene products that are likely involved in DNA synthesis. These include ORF 3 which encodes DNA polymerase, ORF 8 which encodes a SWI/SNF helicase-related protein, ORF 10 encodes a protein showing homology to recA, and ORF 13 encodes a dnaZX-like ORF.

In E. coli, RapA encodes an RNA polymerase (RNAP)-associated protein with ATPase activity and which is a homolog of the eukaryotic SWI/SNF family, a set of proteins whose members are involved are involved in transcription activation, nucleosome remodeling, and DNA repair. RapA forms a stable complex with RNAP,

as if it were a subunit of RNAP and it is possible that the ORF 8 product behaves similarly or in a dominant-negative fashion to inhibit the activity of RapA. Mutation of the essential *E. coli* dnaZX results in a block in DNA chain elongation during replication (Maki et al., 1988). The dnaZX gene has only one open reading frame for a 71-kDa polypeptide from which the two distinct DNA polymerase III holoenzyme subunits, tau (71 kDa) and gamma (47 kDa), are produced. The tau subunit is the precursor of the gamma subunit, and the gamma subunit is produced by a -1 frameshift causing early termination of translation (Tsuchihashi et al., 1990). These proteins show single-strand DNA binding properties that is ATPase (and dATPase) dependent and are thought to increasing the processivity of the core DNA polymerase enzyme (Lee et al., 1987).

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There are several Dp-1 ORFs which encode proteins predicted to play a role in cellular metabolic pathways. These include polypeptides involved in coenzyme PQQ synthesis (ORFs 20, 29, 38). Pyrrolo-quinoline quinone (PQQ) is the non-covalently bound prosthetic group of many quinoproteins catalysing reactions in the periplasm of Gram-negative bacteria. Most of these involve the oxidation of alcohols or aldose sugars. Interestingly, ORFs 20, 29, and 30 also show homology to the exoenzyme S regulon (Frank, 1997). Proteins encoded by the *P. aeruginosa* exoenzyme S regulon may be involved in a contact-mediated translocation mechanism to transfer anti-host factors directly into eukaryotic cells disrupting eukaryotic signal transduction through ADP-ribosylation (Frank, 1997).

There is also a protein with similarity to GTP cyclohydrolase I (ORF 21) and ORF 41 which shows homology to dUTPase (Table 31). GTP cyclohydrolase I is an enzyme that catalyzes the first reaction in the pathway for the biosynthesis of the pteridine, a cofactor of the monooxygenases of the aromatic amino acids. Disruption of the homologous gene in *Saccharomyces cerevisiae* leads to a recessive conditional lethality due to folinic acid auxotrophy, that can be complemented with the mammalian or bacterial GTP cyclohydrolase I enzymes (Nardese et al., 1996; Mancini et al., 1999).

ORF 16 shows high homology to autolysin. This region of the phage sequence was previously reported (Sheehan et al., 1997) and encompasses ~ 4 kbp of our sequence. The sequence published by (Sheehan et al., 1997) is shown in Table 32.

Thus, the present invention provides a nucleic acid sequence obtained from Streptococcus bacteriophage Dp-1 comprising at least a portion of a phage Dp-1 ORF; preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 013 encodes a

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protein with homology to the gamma subunit of DNA polymerase (dnaX gene). This protein may act in a dominant-negative fashion to sequester the host DNA polymerase for its own replication, thus inhibiting host DNA replication. The dnaX gene product is essential for *E. coli* replication (Kodaira et al., 1983).

In certain preferred embodiments of the present invention, the bacterial target of a bacteriophage inhibitor ORF product, e.g., an inhibitory protein or polypeptide, is encoded by a *Streptococcus* nucleic acid coding sequence from a host bacterium for bacteriophage Dp-1. As above, possible target sequences are described herein by reference to sequence source sites. The sequence encoding the target preferably corresponds to a *Streptococcus* nucleic acid sequence available from The Institute for Genomic Research (TIGR), or available from GenBank or other public database. The TIGR *Streptococcus* sequences are publicly available at The Institute for Genomics Research at URL: http://www.tigr.org

The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a Streptococcus pneumoniae coding sequences corresponding to a sequence listed in Table 33 herein. Sequences for other Streptococcal species are also available from TIGR and./or from GenBank. The listing in Table 33 describes Streptococcus sequences currently deposited in GenBank. Again, for the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage Dp-1 host Streptococcus sp. genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

In the various aspects of this invention involving Dp-1 sequences, preferably the sequence is preferably not contained in the sequence described in Sheehan et al., 1997 (Table 32).

## Validating Identified Inhibitory Phage ORFs

A fifth step involves validating the identified phage inhibitor ORF by independent methods, and delineating further possible smaller segments of the ORFs

that have inhibitory activity. Several methods exist to validate the role of the identified ORF as an inhibitor ORF.

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One example utilizes the creation of a mutant variant of the phage ORF in which the candidate ORF carries a partial or complete loss-of-function mutation that is measurable as compared with the non-mutant ORF. Comparison of the effects of expression of the loss of function mutant with the normal ORF provides confirmation of the identification of an inhibitor ORF where the loss-of-function mutant provides a measurably lower level of inhibition, preferably no inhibition. The loss of function may be conditional, e.g., temperature sensitive.

Once validation of the inhibitor ORF is achieved, a bi-directional deletion analysis can be carried out using the same experimental system to identify the minimal polypeptide segment that has inhibitor activity. This may be carried out by a variety of means, e.g., by exonuclease or PCR methodologies, and is used to determine if a relatively small segment of the ORF (i.e., the product of the ORF) still possesses inhibitory activity when isolated away from its native sequence. If so, a portion of the ORF encoding this "active portion" can be used as a template for the synthesis of novel anti-microbial agents and further allowing derivation of the peptide sequence, e.g., using modified peptides and/or peptidomimetics.

In creation of certain peptidomimetics, the peptide backbone is transformed into a carbon-based hydrophobic structure that can retain inhibitor activity against the bacterium. This is done by standard medicinal chemistry methods, typically monitored by measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These mimetics can also represent lead compounds for the development of novel antibiotics.

Recently, a major effort has been undertaken by the pharmaceutical industry and their biotechnology partners for the sequencing of bacterial pathogen genomes. The rationale is that the systematic sequencing of the genome will identify all of the bacterial proteins and therefore this proteome will be the target for designing novel inhibitor antibiotics. Although systematic, this approach has several major problems. The first is that analysis of primary amino acid sequences of bacterial proteins does not immediately reveal which protein will be essential for viability of the bacterium, and target validation is thus a major issue. The second problem is one of redundancy, as several biochemical pathways are either structurally duplicated in bacteria (different isoforms of the same enzyme), or functionally duplicated by the presence of salvage pathways in the event of a metabolic block in one pathway (different nutritional conditions). The third is that even a valid target may not be structurally or

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functionally amenable to inhibition by small molecules because of inaccessibility (sequestration of target).

Therefore, there is considerable interest within the pharmaceutical and biotechnology industry in identifying key targets for drug discovery amongst the mass of novel targets generated by large-scale genomic sequencing projects.

On the other hand, and underscoring the instant invention, the phages herein described have, over millions of years, evolved specific mechanisms to target such key biochemical pathways and proteins. In the few cases where inhibition by phages has been elucidated (e.g., see ref. 3), such bacterial targets are invariably rate-limiting in their respective biochemical pathways, are not redundant, and/or are readily accessible for inhibition by the phage (or by another inhibitory compound). Therefore, the sixth step of this invention involves identifying the host biochemical pathways and proteins that are targeted by the phage inhibitory mechanisms.

# 15 <u>Identifying, Validating, and Characterizing Bacterial Host Target Proteins and</u> <u>Affected Pathways</u>

A rationale for this step is that the inhibitor ORF product from the phage physically interacts with and/or modifies certain microbial host components to block their function. Exemplary approaches which can be used to identify the host bacterial pathways and proteins that interact with, and preferably also are inhibited by, phage ORF product(s) are described below.

One approach is a genetic screen to determine physiological protein:protein interaction, for example, using a yeast two hybrid system. In this assay, the phage ORF is fused to the carboxyl terminus of the yeast Gal4 activation domain II (amino acids 768-881) to create a bait vector. A cDNA library of cloned S. aureus sequences which have been engineered into a plasmid where the S. aureus sequences are fused to the DNA binding domain of Gal4 is also generated. These plasmids are introduced alone, or in combination, into yeast strain Y190 - previously engineered with chromosomally integrated copies of the E. coli lacZ and the selectable HIS3 genes, both under Gal4 regulation (Durfee, T., Becherer, K., Chen, P.-L., Yeh, S.-H., Yang, Y., Kilburn, A.E., Lee, W.-H., and Elledge, S.J. (1993). Genes & Dev. 7, 555-569). If the two proteins expressed in yeast interact, the resulting complex will activate transcription from promoters containing Gal4 binding sites. A lacZ and His3 gene, each driven by a promoter containing Gal4 binding sites, have been integrated into the. genome of the host yeast system used for measuring protein-protein interactions. Such a system provides a physiological environment in which to detect potential protein interactions. This system has been extensively used to identify novel protein-protein

interaction partners and to map the sites required for interaction (for example, to identify interacting partners of translation factors (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). *Mol & Cell Biology* 18, 2697-2711), transcription factors (Katagiri, T., Saito, H., Shinohara, A., Ogawa, H., Kamada, N., Nakamura, Y., and Miki, Y. (1998). Genes, *Chromosomes & Cancer* 21, 217-222), and proteins involved in signal transduction (Endo, T.A., Masuhara, M., Yokouchi, M., Suzuki, R., Sakamoto, H., Mitsui, K., Matsumoto, A., Tanimura, S., Ohtsubo, M., Misawa, H., Miyazaki, T., Leonor N., Taniguchi, T., Fujita, T., Kanakura, Y., Komiya, S., and Yoshimura, A. *Nature*. 387, 921-924). This approach has also been used in many published reports to identify interaction between mammalian viral and mammalian cell proteins.

For example, the non-structural protein NS1 of parvovirus is essential for viral DNA amplification and gene expression and is also the major cytopathic effector of these viruses. A yeast two-hybrid screen with NS1 identified a novel cellular protein of unknown function that interacts with NS-1, called SGT, for small glutamine-rich tetratricopeptide repeat (TPR)-containing protein (Cziepluch C. Kordes E. Poirey R. Grewenig A. Rommelaere, J, and Jauniaux JC. (1998) J Virol. 72, 4149-4156). In another screen, the adenovirus E3 protein was recently shown to interact with a novel tumor necrosis factor alpha-inducible protein and to modulate some of the activities of E3 (Li Y. Kang J. and Horwitz M.S. (1998). Mol & Cell Biol. 18, 1601-1610). In yet another recent screen, the herpes simplex virus 1 alpha regulatory protein ICP0 was found to interact with (and stabilize) the cell cycle regulator cyclin D3 (Kawaguchi Y. Van Sant C. and Roizman B. (1997). J Virol. 71,7328-7336).

Another two-hybrid system for identifying protein:protein interactions is commercially available from STRATEGENE<sup>TM</sup> as the CYTO-TRAP<sup>TM</sup> system (Chang et al., *Strategies Newsletter* 11(3), 65-68 (1998)(from Stratagene)). The system is a yeast-based method for detecting protein:protein interactions *in vivo*, using activation of the Ras signal transduction cascade by localizing a signal pathway component, human Sos (hSos), to its activation site in the yeast plasma membrane. The system uses a temperature-sensitive *Saccharomyces cerevisiae* mutant, strain cdc25H, which contains a point mutation at amino acid residue 1328 of the cdc25 gene. This gene encodes a guanyl nucleotide exchange factor which binds and activates Ras, leading to cell growth. The mutation in the cdc25 gene prevents host growth at 37°C, but at a permissive temperature of 25°C, growth is normal. The system utilizes the ability of (hSos) to complement the cdc25 defect and activate the yeast Ras signaling pathway. Once (hSos) is expressed and localized to the plasma membrane, the cdc25H yeast strain grows at 37°C. Localizing hSos to the plasma

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membrane occurs through a protein:protein interaction. A protein of interest, or bait, is expressed as a fusion protein with hSos. The library, or target proteins are expressed with the myristylation membrane-localization signal. The yeast cells are then incubated under restrictive conditions (37°C). If the bait and the target protein interact, the hSos protein is recruited to the membrane, activating the Ras signaling pathway and allowing the cdc25H yeast strain to grow at the restrictive temperature.

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The protein targets of phage inhibitory ORFs can also be identified using bacterial genetic screens. One approach involves the overexpression of a phage inhibitory protein in mutagenized bacterial host species, followed by plating the cells and searching for colonies that can survive the antimicrobial activity of the inhibitory ORF. These colonies are then grown, their DNA extracted, and cloned into an expression vector that contains a replicon of a different incompatibility group from the plasmid expressing the original ORF. This library is then introduced into a wild-type host bacterium in conjunction with an expression vector driving synthesis of the phage ORF, followed by selection for surviving bacteria. Thus, bacterial DNA fragments from the survivors presumably contain a DNA fragment from the original mutagenized host bacterial genome that can protect the cell from the antimicrobial activity of the inhibitory phage ORF. This fragment can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach enables one to determine the targets and pathways that are affected by the killing function.

A second approach is based on identifying protein:protein interactions between the phage ORF product and bacterial S. aureus, e.g., proteins using a biochemical approach based, for example, on affinity chromatography. This approach has been used, for example, to identify interactions between lambda phage proteins and proteins from their E. coli host (Sopta, M., Carthew, R.W., and Greenblatt, J. (1985) J. Biol. Chem. 260, 10353-10369). The phage ORF is fused to a peptide tag (e.g. glutathione-S-transferase ("GST"), 6xHIS, ("HIS") and/or calmodulin binding protein ("CPB")) within a commercially available plasmid vector that directs high level expression on induction of a suitably responsive promoter driving the fusion's expression. The translated fusion protein is expressed in E. coli, purified, and immobilized on a solid phase matrix via, for example the tag. Total cell extracts from the host bacterium, e.g., S. aureus, are then passed through the affinity matrix containing the immobilized phage ORF fusion protein; host proteins retained on the column are then eluted under different conditions of ionic strength, pH, detergents etc., and characterized by gel electrophoresis and other techniques. Appropriate controls are run to guard against nonspecific binding to the resin. Target proteins thus

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recovered should be enriched for the phage protein/peptide of interest and are subsequently electrophoretically or otherwise separated, purified, sequenced, or biochemically analyzed. Usually sequencing entails individual digestion of the proteins to completion with a protease (e.g.-trypsin), followed by molecular mass and amino acid composition and sequence determination using, for example, mass spectrometry, e.g., by MALDI-TOF technology (Qin, J., Fenyo, D., Zhao, Y., Hall, W.W., Chao, D.M., Wilson, C.J., Young, R.A. and Chait, B.T. (1997). Anal. Chem. 69, 3995-4001).

The sequence of the individual peptides from a single protein are then analyzed by the bioinformatics approach described above to identify the *S. aureus* protein interacting with the phage ORF. This analysis is performed by a computer search of the *S. aureus* genome for an identified sequence. Alternatively, all tryptic peptide fragments of the *S. aureus* genome can be predicted by computer software, and the molecular mass of such fragments compared to the molecular mass of the peptides obtained from each interacting protein eluted from the affinity matrix. The responsible gene sequence can be obtained, for example by using synthetic degenerate nucleic acid sequences to pull out the corresponding homologous bacterial sequence. Alternatively, antibodies can be generated against the peptide and used to isolate nascent peptide/mRNA transcript complexes, from which the mRNA can be reverse transcribed, cloned, and further characterized using the procedures discussed herein.

A variety of other binding assay methods are known in the art and can be used to identify interactions between phage proteins and bacterial proteins or other bacterial cell components. Such methods that allow or provide identification of the bacterial component can be used in this invention for identifying putative targets.

Validation of the interaction between the phage ORF product and the bacterial proteins or other components can be obtained by a second independent assay (e.g., co-immunoprecipitation or protein-protein crosslinking experiments (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). Mol & Cell Biology 18, 2697-2711; Brown, S. and Blumenthal, T. (1976). Proc. Natl. Acad. Sci. USA 73, 1131-1135)).

Finally, the essential nature of the identified bacterial proteins is preferably determined genetically by creating a constitutive or inducible partial or complete loss-of-function mutation in the gene encoding the identified interacting bacterial protein. This mutant is then tested for bacterial survival and replication.

The protein target of the phage inhibitor function can also be identified using a genetic approach. Two exemplary approaches will be delineated here. The first approach involves the overexpression of a predetermined phage inhibitor protein in mutagenized host bacteria, e.g., S. aureus, followed by plating the cells and searching

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for colonies that can survive the inhibitor. These colonies will then be grown, their DNA extracted and cloned into an expression vector that contains a replicon of a different incompatibility group, and preferably having a different selectible marker than the plasmid expressing the phage inhibitor. Thus, host DNA fragments from the mutant that can protect the cell from phage ORF inhibition can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach allows rapid determination of the targets and pathways that are affected by the inhibitor.

Alternatively, the bacterial targets can be determined in the absence of selecting for mutations using an approach known as "multicopy suppression". In this approach, the DNA from the wild type host is cloned into an expression vector that can coexist, as previously described, with one containing a predetermined phage inhibitor. Those plasmids that contain host DNA fragments and genes that protect the host from the phage inhibitor can then be isolated and sequenced to identify putative targets and pathways in the host bacteria.

Regardless of the specific mode of identification, screening assays may additionally utilize gene fusions to specific "reporter genes" to identify a bacterial gene(s) whose expression is affected when the host target pathway is affected by the phage inhibitor. Such gene fusions can be used to search a number of small molecule compounds for inhibitors that may affect this pathway and thus cause cell inhibition. This approach will allow the screening of a large number of molecules on petri dishes or 96-well format by monitoring for a simple color change in the bacterial colonies. In this manner, we can validate host targets and classes of compounds for further study and clinical development. These inhibitors also represent lead compounds for the development of other antibiotics.

Bioinformatics and comparative genomics are preferably then applied to the identified bacterial gene products to predict biochemical function. The biochemical activity of the protein can be verified *in vitro* in cell free assays or *in vivo* in intact cells. *In vitro* biochemical assays utilizing cell-free extracts or purified protein are established as a basis for the screening and development of inhibitors.

These inhibitors, preferably small molecule inhibitors, may comprise peptides, antibodies, products from natural sources such as fungal or plant extracts or small molecule organic compounds. In general, small molecule organic compounds are preferred. These compounds may, for example, be identified within large compound libraries, including combinatorial libraries. For example, a plurality of compounds, preferably a large number of compounds can be screened to determine whether any of the compounds binds or otherwise disrupts or inhibits the identified bacterial target.

Compounds identified as having any of these activities can then be evaluated further in cell culture and/or animal model systems to determine the pharmacological properties of the compound, including the specific anti-microbial ability of the compound.

For mixtures of natural products, including crude preparations, once a preparation or fraction of a preparation is shown the have an anti-microbial activity, the active substance can be isolated and identified using techniques well known in the art, if the compound is not already available in a purified form.

Identified compounds possessing anti-microbial activity and similar compounds having structural similarity can be further evaluated and, if necessary, derivatized according to synthesis and/or modification methods available in the art selected as appropriate for the particular starting molecule.

# Derivatization of identified anti-microbials

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In cases where the identified anti-microbials above might represent peptidal compunds, the *in vivo* effectiveness of such compounds may be advantageously enhanced by chemical modification using the natural polypeptide as a starting point and incorporating changes that provide advantages for use, for example, increased stability to proteolytic degradation, reduced antigenicity, improved tissue penetration, and/or improved delivery characteristics.

In addition to active modifications and derivative creations, it can also be useful to provide inactive modifications or derivatives for use as negative controls or introduction of immunologic tolerance. For example, a biologically inactive derivative which has essentially the same epitopes as the corresponding natural antimicrobial can be used to induce immunological tolerance in a patient being treated. The induction of tolerance can then allow uninterrupted treatment with the active anti-microbial to continue for a significantly longer period of time.

Modified anti-microbial polypeptides and derivatives can be produced using a number of different types of modifications to the amino acid chain. Many such methods are known to those skilled in the art. The changes can include, for example, reduction of the size of the molecule, and/or the modification of the amino acid sequence of the molecule. In addition, a variety of different chemical modifications of the naturally occurring polypeptide can be used, either with or without modifications to the amino acid sequence or size of the molecule. Such chemical modifications can, for example, include the incorporation of modified or non-natural amino acids or non-amino acid moieties during synthesis of the peptide chain, or the post-synthesis modification of incorporated chain moieties.

The oligopeptides of this invention can be synthesized chemically or through an appropriate gene expression system. Synthetic peptides can include both naturally occurring amino acids and laboratory synthesized, modified amino acids.

Also provided herein are functional derivatives of anti-microbial proteins or polypeptides. By "functional derivative" is meant a "chemical derivative," "fragment," "variant," "chimera," or "hybrid" of the polypeptide or protein, which terms are defined below. A functional derivative retains at least a portion of the function of the protein, for example reactivity with a specific antibody, enzymatic activity or binding activity.

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A "chemical derivative" of the complex contains additional chemical moieties not normally a part of the protein or peptide. Such moieties may improve the molecule's solubility, absorption, biological half-life, and the like. The moieties may alternatively decrease the toxicity of the molecule, eliminate or attenuate any undesirable side effect of the molecule, and the like. Moieties capable of mediating such effects are disclosed in Alfonso and Gennaro (1995). Procedures for coupling such moieties to a molecule are well known in the art. Covalent modifications of the protein or peptides are included within the scope of this invention. Such modifications may be introduced into the molecule by reacting targeted amino acid residues of the peptide with an organic derivatizing agent that is capable of reacting with selected side chains or terminal residues, as described below.

Cysteinyl residues most commonly are reacted with alpha-haloacetates (and corresponding amines), such as chloroacetic acid or chloroacetamide, to give carboxymethyl or carboxyamidomethyl derivatives. Cysteinyl residues also are derivatized by reaction with bromotrifluoroacetone, chloroacetyl phosphate, N-alkylmaleimides, 3-nitro-2-pyridyl disulfide, methyl 2-pyridyl disulfide, p-chloromercuribenzoate, 2-chloromercuri-4-nitrophenol, or chloro-7-nitrobenzo-2-oxa-1,3-diazole.

Histidyl residues are derivatized by reaction with diethylprocarbonate at pH 5.5-7.0 because this agent is relatively specific for the histidyl side chain. Parabromophenacyl bromide also is useful; the reaction is preferably performed in 0.1 M sodium cacodylate at pH 6.0.

Lysinyl and amino terminal residues are reacted with succinic or other carboxylic acid anhydrides. Derivatization with these agents has the effect of reversing the charge of the lysinyl residues. Other suitable reagents for derivatizing primary amine- containing residues include imidoesters such as methyl picolinimidate; pyridoxal phosphate; pyridoxal; chloroborohydride;

trinitrobenzenesulfonic acid; O-methylisourea; 2,4 pentanedione; and transaminasecatalyzed reaction with glyoxylate.

Arginyl residues are modified by reaction with one or several conventional reagents, among them phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, and ninhydrin. Derivatization of arginine residues requires that the reaction be performed in alkaline conditions because of the high pK<sub>2</sub> of the guanidine functional group. Furthermore, these reagents may react with the groups of lysine as well as the arginine alpha-amino group.

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Tyrosyl residues are well-known targets of modification for introduction of spectral labels by reaction with aromatic diazonium compounds or tetranitromethane. Most commonly, N-acetylimidizol and tetranitromethane are used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively.

Carboxyl side groups (aspartyl or glutamyl) are selectively modified by reaction carbodiimide (R'-N-C-N-R') such as 1-cyclohexyl-3-(2-morpholinyl(4-ethyl) carbodiimide or 1-ethyl-3-(4-azonia-4,4-dimethylpentyl) carbodiimide. Furthermore, aspartyl and glutamyl residues are converted to asparaginyl and glutaminyl residues by reaction with ammonium ions.

Glutaminyl and asparaginyl residues are frequently deamidated to the corresponding glutamyl and aspartyl residues. Alternatively, these residues are deamidated under mildly acidic conditions. Either form of these residues falls within the scope of this invention.

Derivatization with bifunctional agents is useful, for example, for cross-linking component peptides to each other or the complex to a water-insoluble support matrix or to other macromolecular carriers. Commonly used cross-linking agents include, for example, 1,1-bis (diazoacetyl)-2-phenylethane, glutaraldehyde, N-hydroxysuccinimide esters, for example, esters with 4-azidosalicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis(succinimidylpropionate), and bifunctional maleimides such as bis-N-maleimido-1,8-octane. Derivatizing agents such as methyl-3-[p-azidophenyl) dithiolpropioimidate yield photoactivatable intermediates that are capable of forming crosslinks in the presence of light. Alternatively, reactive water-insoluble matrices such as cyanogen bromide-activated carbohydrates and the reactive substrates described in U.S. Patent Nos. 3,969,287; 3,691,016; 4,195,128; 4,247,642; 4,229,537; and 4,330,440 are employed for protein immobilization.

Other modifications include hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the alpha-amino groups of lysine, arginine, and histidine side chains (Creighton, T.E.,

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Proteins: Structure and Molecular Properties, W.H. Freeman & Co., San Francisco, pp. 79-86 (1983)), acetylation of the N-terminal amine, and, in some instances, amidation of the C-terminal carboxyl groups.

Such derivatized moieties may improve the stability, solubility, absorption, biological half life, and the like. The moieties may alternatively eliminate or attenuate any undesirable side effect of the protein complex. Moieties capable of mediating such effects are disclosed, for example, in Alfonso and Gennaro (1995).

The term "fragment" is used to indicate a polypeptide derived from the amino acid sequence of the protein or polypeptide having a length less than the full-length polypeptide from which it has been derived. Such a fragment may, for example, be produced by proteolytic cleavage of the full-length protein. Preferably, the fragment is obtained recombinantly by appropriately modifying the DNA sequence encoding the proteins to delete one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

Another functional derivative intended to be within the scope of the present invention is a "variant" polypeptide that either lacks one or more amino acids or contains additional or substituted amino acids relative to the native polypeptide. The variant may be derived from a naturally occurring polypeptide by appropriately modifying the protein DNA coding sequence to add, remove, and/or to modify codons for one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

A functional derivative of a protein or polypeptide with deleted, inserted and/or substituted amino acid residues may be prepared using standard techniques well-known to those of ordinary skill in the art. For example, the modified components of the functional derivatives may be produced using site-directed mutagenesis techniques (as exemplified by Adelman et al., 1983, *DNA* 2:183; Sambrook et al., 1989) wherein nucleotides in the DNA coding sequence are modified such that a modified coding sequence is produced, and thereafter expressing this recombinant DNA in a prokaryotic or eukaryotic host cell, using techniques such as those described above. Alternatively, components of functional derivatives of complexes with amino acid deletions, insertions and/or substitutions may be conveniently prepared by direct chemical synthesis, using methods well-known in the art.

Insofar as other anti-microbial inhibitor compounds identified by the invention described herein may not be peptidal in nature, other chemical techniques exist to allow their suitable modification, as well, and according the desirable principles discussed above.

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## Administration and Pharmaceutical Compositions

For the therapeutic and prophylactic treatment of infection, the preferred method of preparation or administration of anti-microbial compounds will generally vary depending on the precise identity and nature of the anti-microbial being delivered. Thus, those skilled in the art will understand that administration methods known in the art will also be appropriate for the compounds of this invention.

The particularly desired anti-microbial can be administered to a patient either by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s). In treating an infection, a therapeutically effective amount of an agent or agents is administered. A therapeutically effective dose refers to that amount of the compound that results in amelioration of one or more symptoms of bacterial infection and/or a prolongation of patient survival or patient comfort.

Toxicity, therapeutic and prophylactic efficacy of anti-microbials can be determined by standard pharmaceutical procedures in cell cultures and/or experimental organisms such as animals, e.g., for determining the LD<sub>50</sub> (the dose lethal to 50% of the population) and the ED<sub>50</sub> (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD<sub>50</sub>/ED<sub>50</sub>. Compounds that exhibit large therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED<sub>50</sub> with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized.

For any compound identified and used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. Such information can be used to more accurately determine useful doses in organisms such as plants and animals, preferably mammals, and most preferably humans. Levels in plasma may be measured, for example, by HPLC or other means appropriate for detection of the particular compound.

The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition (see *e.g.* Fingl et. al., in The Pharmacological Basis of Therapeutics, 1975, Ch. 1 p.1).

It should be noted that the attending physician would know how and when to terminate, interrupt, or adjust administration due to toxicity, organ dysfunction, or other systemic malady. Conversely, the attending physician would also know to adjust treatment to higher levels if the clinical response were not adequate (precluding

toxicity). The magnitude of an administered dose in the management of the disorder of interest will vary with the severity of the condition to be treated and the route of administration. The severity of the condition may, for example, be evaluated, in part, by standard prognostic evaluation methods. Further, the dose and perhaps dose frequency, will also vary according to the age, body weight, and response of the individual patient. A program comparable to that discussed above also may be used in veterinary or phyto medicine.

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Depending on the specific infection target being treated and the method selected, such agents may be formulated and administered systemically or locally, i.e., topically. Techniques for formulation and administration may be found in Alfonso and Gennaro (1995). Suitable routes may include, for example, oral, rectal, transdermal, vaginal, transmucosal, intestinal, parenteral, intramuscular, subcutaneous, or intramedullary injections, as well as intrathecal, intravenous, or intraperitoneal injections.

For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks' solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

Use of pharmaceutically acceptable carriers to formulate identified antimicrobials of the present invention into dosages suitable for systemic administration is within the scope of the invention. With proper choice of carrier and suitable manufacturing practice, the compositions of the present invention, in particular those formulated as solutions, may be administered parenterally, such as by intravenous injection. Appropriate compounds can be formulated readily using pharmaceutically acceptable carriers well known in the art into dosages suitable for oral administration. Such carriers enable the compounds of the invention to be formulated as tablets, pills, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated.

Agents intended to be administered intracellularly may be administered using techniques well known to those of ordinary skill in the art. For example, such agents may be encapsulated into liposomes, then administered as described above.

Liposomes are spherical lipid bilayers with aqueous interiors. All molecules present in an aqueous solution at the time of liposome formation are incorporated into the aqueous interior. The liposomal contents are both protected from the external microenvironment and, because liposomes fuse with cell membranes, are efficiently

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delivered into the cell cytoplasm. Additionally, due to their hydrophobicity, small organic molecules may be directly administered intracellularly.

Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to achieve the intended purpose. Determination of the effective amounts is well within the capability of those skilled in the art.

In addition to the active ingredients, these pharmaceutical compositions may contain suitable pharmaceutically acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. The preparations formulated for oral administration may be in the form of tablets, dragees, capsules, or solutions, including those formulated for delayed release or only to be released when the pharmaceutical reaches the small or large intestine.

The pharmaceutical compositions of the present invention may be manufactured in a manner that is itself known, e.g., by means of conventional mixing, dissolving, granulating, dragee-making, levitating, emulsifying, encapsulating, entrapping or lyophilizing processes.

Pharmaceutical formulations for parenteral administration include aqueous solutions of the active anti-microbial compounds in water-soluble form.

Alternatively, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions.

Pharmaceutical preparations for oral use can be obtained by combining the active compounds with solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

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Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures.

Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added.

The above methodologies may be employed either actively or prophylactically against an infection of interest.

#### Computer-related Aspects and Embodiments

In addition to the provision of compounds as chemical entities, nucleotide sequences, or fragments thereof at least 95%, preferably at least 97%, more preferably at least 99%, and most preferably at least 99.9% identical to phage inhibitor sequences can also be provided in a variety of additional media to facilitate various uses.

Thus, as used in this section, "provided" refers to an article of manufacture, rather than an actual nucleic acid molecule, which contains a nucleotide sequence of the present invention; e.g., a nucleotide sequence of an exemplary bacteriophage or a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of an unsequenced phage listed in Table 1, preferably of bacteriophage 77 (S. aureus host) or bacteriophage 3A (S. aureus host) or bacteriophage 96 (S. aureus host). Such an article provides a large portion of the particular bacteriophage genome or bacterial gene and parts thereof (e.g., a bacteriophage open reading frame (ORF)) in a form which allows a skilled artisan to examine and/or analyze the sequence using means not directly applicable to examining the actual genome or gene or subset thereof as it exists in nature or in purified form as a chemical entity.

In one application of this aspect, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer

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readable media" refers to any medium that can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories, such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create an article of manufacture which includes one or more computer readable media having recorded thereon a nucleotide sequence or sequences of the present invention. Likewise, it will be clear to those of skill how additional computer readable media that may be developed also can be used to create analogous manufactures having recorded thereon a nucleotide sequence of the present invention.

As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can, for example, be presented in a word processing test file, formatted in commercially available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g., text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. Thus, by providing in computer readable form a nucleotide sequence of an unsequenced bacteriophage, such as an exemplary bacteriophage listed in Table 1 or of a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of bacteriophage 77 (S. aureus host) or bacteriophage 3A (S. aureus host) bacteriophage

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96 (S. aureus host), bacteriophage 44AHJD (S. aureus host), bacteriophage Dp-1 (Streptococcus pneumoniae host), or bacteriophage 182 (Enterococcus host) the present invention enables the skilled artisan to routinely access the provided sequence information for a wide variety of purposes.

Those skilled in the art understand that software can implement a variety of different search or analysis software which implement sequence search and analysis algorithms, e.g., the BLAST (Altschul et al., J. Mol. Biol. 215:403410 (1990) and BLAZE (Brutlag et al., Comp. Chem 17:203-207 (1993)) search algorithms. For example, such search algorithms can be implemented on a Sybase system and used to identify open reading frames (ORFs) within the bacteriophage genome which contain homology to ORFs or proteins from other viruses, e.g., other bacteriophage, and other organisms, e.g., the host bacterium. Among the ORFs discussed herein are protein encoding fragments of the bacteriophage genomes which encode bacteria-inhibiting proteins or fragments.

The present invention further provides systems, particularly computer-based systems, which contain the sequence information described. Such systems are designed to identify, among other things, useful fragments of the bacteriophage genomes.

As used herein, "a computer-based system" refers to the hardware, software, and data storage media used to analyze the nucleotide sequence information of the present invention. The minimum hardware of the computer-based systems of the present invention comprises a central processing unit (CPU), input device, output device, and data storage medium or media. A skilled artisan will readily recognize that any of the currently available general purpose computer-based system are suitable for use in the present invention, as well as a variety of different specialized or dedicated computer-based systems.

As stated above, the computer-based systems of the present invention comprise data storage media having stored therein a nucleotide sequence of the present invention and the necessary hardware and software for supporting and implementing a search and/or analysis program.

As used herein, "data storage media" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

As used herein, "search program" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means.

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Search means are used to identify fragments or regions of the present gnomic sequences which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, MacPattern (EMBL), BLASTN and BLASTX (NCBIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches and/or sequence analyses can be adapted for use in the present computer-based systems.

As used herein in connection with sequence searches and analyses, a "target sequence" can be any DNA or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. Also, the target sequence length is preferably selected to include sequence corresponding to a biologically relevant portion of an encoded product, for example a region which is expected to be conserved across a range of source organisms. Preferably the sequence length of a target polypeptide sequence is from 5-100 amino acids, more preferably 7-50 or 7-100 amino acids, and still more preferably 10-80 or 10-100 amino acids. Preferably the sequence length of a target polynucleotide sequence is from 15-300 nucleotide residues, more preferably from 21-240 or 21-300, and still more preferably 30-150 or 30-300 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length. Likewise, it may be desirable to search and/or analyze longer sequences.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzymatic active sites and signal sequences. Nucleic acid target motifs include, but are not limited to promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

A variety of structural formats for the input and output devices can be used to input and output the information in the computer-based systems of the present invention. A preferred format for an output device ranks fragments of the bacteriophage or bacterial sequences possessing varying degrees of homology to the

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target sequence or target motif. Such presentation provides a skilled artisan with a ranking of sequences which contain various amounts of the target sequence or target motif and identifies the degree of homology contained in the identified fragment.

A variety of comparing methods and/or devices and/or formats can be used to compare a target sequence or target motif with the sequence stored in data storage media to identify sequence fragments of the bacteriophage or bacterium in question. One skilled in the art can readily recognize that any one of the publicly available homology search programs can be used as the search program for the computer-based systems of the present invention. Of course, suitable proprietary systems that may be known to those of skill, or later developed, also may be employed in this regard.

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Figure 6 provides a block diagram of a computer system illustrative of embodiments of this aspect of present invention. The computer system 102 includes a processor 106 connected to a bus 104. Also connected to the bus 104 are a main memory 108 (preferably implemented as random access memory, RAM) and a variety of secondary storage devices 110, such as a hard drive 112 and a removable medium storage device 114. The removable medium storage device114 may represent, for example, a floppy disk drive, a CD-ROM drive, a magnetic tape drive, etc. A removable storage medium 116 (such as a floppy disk, a compact disk, a magnetic tape, etc.) containing control logic and/or data recorded therein may be inserted into the removable medium storage device 114. The computer system 102 includes appropriate software for reading the control logic and/or the data from the removable medium storage device 114, once it is inserted into the removable medium storage device 114.

A nucleotide sequence of the present invention may be stored in a well-known manner in the main memory 108, any of the secondary storage devices 110, and/or a removable storage medium 116. During execution, software for accessing and processing the sequence (such as search tools, comparing tools, etc.) reside in main memory 108, in accordance with the requirements and operating parameters of the operating system, the hardware system and the software program or programs.

The data storage medium in which the sequence is embodied and the central processor need not be part of a single stand-alone computer, but may be separated so long as data transfer can occur. For example, the processor or processors being utilized for a search or analysis can be part of one general purpose computer, and the data storage medium can be part of a second general purpose computer connected to a network, or the data storage medium can be part of a network server. As another example the data storage medium can be part of a computer system or network accessible over telephone lines or other remote connection method.

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#### **EXAMPLES**

## Example 1. Growth of Staph A bacteriophage 77 and purification of genomic DNA.

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The Staphylococcus aureus propagating strain (PS 77; ATCC #27699) was used as a host to propagate its respective phage 77 (ATCC # 27699-B1). Two rounds of plaque purification of phage 77 were performed on soft agar essentially as described in Sambrook et al (1989). Briefly, the PS 77 strain was grown overnight at 37°C in Nutrient broth [NB: 0.3% Bacto beef extract, 0.5% Bacto peptone (Difco Laboratories) and 0.5% NaCl (w/v)]. The culture was then diluted 20x in NB and incubated at 37°C until the OD<sub>540</sub>= .2 (early log phase) with constant agitation. In order to obtain single plaques, phage 77 was subjected to 10-fold serial dilutions using phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 µl of each dilution was used to infect 0.5 ml of the cell suspension in the presence of 400 µg/ml CaCl<sub>2</sub>. After incubation of 15 min at room temperature (RT), 2 ml of melted soft agar kept at 45°C (NB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm nutrient agar plates (0.3% Bacto Beef extract, 0.5% Bacto peptone, 0.5% NaCl and 1.5% Bacto agar (w/v)). After overnight incubation at 30°C, a single plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at 20°C, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 30°C, a single plaque was isolated and used as a stock.

The propagation procedure for bacteriophage 77 was modified from the agar layer method of Swanstörm and Adams (1951). Briefly, the PS 77 strain was grown to stationary phase overnight at 37°C in Nutrient broth. The culture was then diluted twenty-fold in NB and incubated at 37°C until the OD<sub>540</sub>= .2. The suspension (15x10<sup>7</sup> Bacteria) was then mixed with 15x10<sup>5</sup> plaque forming units (pfu) to give a ratio of 100-bacteria/phage particle in the presence of 400 μg/ml of CaCl<sub>2</sub>. After incubation for 15 min at 20°C, 7.5 ml of melted soft agar (NB plus 0.6% agar) were added to the mixture and poured onto the surface of 150 mm nutrient agar plates and incubated 16 hrs at 30°C. To collect the phage plate lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide followed by shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 RPM (2,830xg) in a JA-10 rotor-(Beckman) and the supernatant fluid (lysate) was collected and subjected to a treatment with 10 μg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) PEG 8000 and

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0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS 55 rotor centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 rpm (67,000xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000xg) for 24 h at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl<sub>2</sub>. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 mg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris pH 8.0, 1mM EDTA).

#### Example 2. DNA sequencing of Bacteriophage 77 genome

Four micrograms of phage 77 DNA was diluted in 200 μl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator<sup>TM</sup>, Fisher Scientific). Samples were sonicated under an amplitude of 3 μm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 μl of 1 mM Tris (pH 8.5).

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl<sub>2</sub>, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of Klenow large fragment (New England Biolabs) for 15 min at room—temperature. The reaction was stopped by two phenol/chloroform extractions and the

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DNA was precipitated with ethanol and the final DNA pellet was resuspended in 20  $\mu$ l of  $H_2O$ .

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs)-treated pKS II+ vector (Stratagene). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 μl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 μl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook et al., 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μl LB and 100 μg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS II+ vector. PCR amplification of foreign insert was performed in a 15 μl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 μM primer, 187.5 μM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 57°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using OIAprep<sup>TM</sup> spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye<sup>TM</sup> primer or ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems). To ensure co-linearity of the sequence data and the genome, all regions of phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit.

## Example 3. Bioinformatic management of primary nucleotide sequence from 35 Phage 77.

Phage 77 sequence contigs were assembled using Sequencher<sup>TM</sup> 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of

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the contigs. Phage DNA was used directly as sequencing template employing ABI prism BIG DYE™ terminator cycle sequencing ready reaction kit. The complete sequence of bacteriophage 77 is shown in Table 2.

A software program was developed and used on the assembled sequence of bacteriophage 77 to identify all putative ORFs larger than 33 codons. Other ORF identification software can also be utilized, preferably programs which allow alternative start codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI (<a href="http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c">http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c</a>) for the bacterial genetic code.

When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons (start and stop codons) is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence.

Sequence homology (BLAST) searches for each ORF are then carried out using an implementation of BLAST programs, although any of a variety of different sequence comparison and matching programs can be utilized as known to those skilled in the art. Downloaded public databases used for sequence analysis include:

- i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
  - ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
  - iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
  - iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
- 30 v) S. aureus NCTC 8325 (ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa);
  - vi) streptococcus pyogenes (ftp://ftp.genome.ou.edu/pub/strep/strep-1k.fa);
  - vii) Streptococcus pneumoniae
  - (ftp://ftp.tigr.org/pub/data/s\_pneumoniae/gsp.contigs.112197.Z);
  - viii) Mycobacterium tuberculosis CSU#9
- 35 (ftp://ftp.tigr.org/pub/data/m\_tuberculosis/TB\_091097.Z) and ix) pseudomonas aeruginosa (http://www.genome.washington.edu/pseudo/data.html).

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The results of the homology searches performed on the ORFs is shown in Table 5.

## Example 4. Subcloning of Bacteriophage 77 ORFs into a Staph A inducible expression system.

The shuttle vector pT0021, in which the firefly luciferase (*lucFF*) expression is controlled by the *ars* (arsenite) promoter/operator (Tauriainen et al., 1997), was modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the heamaglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with *BamH*I, *SaI*I and *Hind*III cloning sites) is: 5'-gatcccggtcgaccaagcttTACCCATACGACGTCCCAGACTACGCCAGCTGA-3' (where upper case letters denote the nucletotide sequence of the HA tag); the antisense strand HA tag sequence (with a *Hind*III cloning site) is:

5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAaagcttggtcgaccgg-3' (where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with BamHI and HindIII. This manipulation resulted in replacement of the lucFF gene by the HA tag. This modified shuttle vector containing the arsenite
 inducible promoter, the arsR gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A.

Each ORF, encoded by Bacteriophage 77, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon was selected for functional analysis for bacterial inhibition. In total, 98 ORFs were selected and screened as detailed below. A list of these is presented in Table 3. Each individual ORF, from initiation codon to last codon (excluding the stop codon), was amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of ORFs, each sense strand primer targets the initiation codon and is preceded by a BamHI restriction site (5'cgggatcc3') and each antisense oligonucleotide targets the pentultimate codon (the one before the stop codon) of the ORF and is preceded by a Sal I restriction site (<sup>5</sup>gcgtcgaccg<sup>3</sup>). The PCR product of each ORF was gel purified and digested with BamHI and SalI. The digested PCR product was then gel purified using the Qiagen kit as described, ligated into BamHI and SalI digested pTHA vector, and used to transform E. coli bacterial strain DH10β(as described above). As a result of this manipulation, the HA tag is set inframe with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones were picked and their insert sizes were confirmed by PCR analysis

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using primers flanking the cloning site. The names and sequences of the primers that were used for the PCR amplification were: HAF:

<sup>5</sup>TATTATCCAAAACTTGAACA<sup>3</sup>; HAR: <sup>5</sup>CGGTGGTATATCCAGTGATT<sup>3</sup>. The sequence integrity of cloned ORFs was verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence could not be achieved by one pass with the sequencing primers, additional internal primers were selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiswirth et al., 1983) was used as a recipient for the expression of recombinant plasmids. Electoporation was performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones was performed on Luria-Broth agar (LB-agar) plates containing 30 µg/ml of kanamycin.

For each ORF introduced in the pTHA plasmid, 3 independent transformants were isolated and used to individually inoculate cultures in 5 ml of TSB containing 15 30µg/ml kanamycin, followed by growth to saturation (16 hrs at 30°C). An aliquot of this stationary phase culture was used to generate a frozen glycerol stock of the transformant ( stored at - 80°C). The remaining culture was used for plasmid DNA extraction. Bacterial cells were harvested by centrifugation at 3000 x g at 22°C for 5 min. The pellet was resuspended in 200 µl 25% sucrose containing 25U/ml of 20 lysostaphin and incubated for 15 min at 37°C. Then, 400µl of alkaline SDS solution (3% SDS, 0.2N NaOH) were added, well mixed and incubated for 7 min at room temperature. After the alkaline SDS treatment, 300µl of ice-cold 3M sodium acetate pH 4.8 were added, and the mix is immediately spun at 13000g for 15 min at room temperature. The supernatant was transferred to a new 1.5 ml conical centrifuge tube 25 and 650µl of isopropanol (stored at room temperature) were added. The mix was then centrifuged at 13,000 x g for 5 min. The supernatant fluid was discarded, the pellet washed with 70% ethanol, and resuspended in 320 µl sterile distilled water.

The presence of individual phage 77 ORF DNA inserts in the plasmid was verified by PCR amplification using 1.5  $\mu$ l transformant miniprep DNA in a PCR with primers flanking the cloning site of ORF in pTHA vector (HAF and HAR). The composition of the PCR reaction and the cycling parameters are identical to those employed for library screening described above.

Example 5. Functional assay for bacterial inhibitory activity of bacteriophage 77

35 ORFs.

The anti-microbial activity of individual phage 77 ORFs was monitored by two growth inhibitory assays, one on solid agar medium, the other in liquid medium.

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In general, Staphylococcus bacteria transformed with expression plasmids containing individual ORFs were grown in normal TSA medium and stored in 19% glycerol. At pre-determined times, arsenite was added to the culture to induce transcription of the phage 77 ORFs cloned immediately downstream from an arsenite-inducible promoter in the pTHA expression plasmid.

The effect of ORF induction on bacterial growth characteristics was then monitored and quantitated. The growth inhibition assay on solid medium was performed by streaking pTHA/ORF containing *S. aureus* transformant onto LB-Kn and TSA-Kn plates containing increasing concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μM). Arsenite is used to induce the expression of cloned DNA in pTHA vector. In parallel, 3 μl of 1/10 and 1/100 dilutions of the frozen cultures of the pTHA/ORF transformants were spotted as single drops onto LB-Kn and TSA-Kn plates containing increasing concentration of sodium arsenite (0; 2.5; 5; and 7.5 μM). The plates were then incubated 16 hrs at 37°C, and the effect of arsenite-induced ORF expression on bacterial growth was monitored and quantitated by comparing the extent to that seen in control plates. As positive controls for growth inhibition,the *holin/lysin* genes of the *Sthaphylococcus aureus* phage Twort (Loessner et al., 1998) was subcloned into the pTHA *ars* inducible vector and used.

For the growth inhibition assay in liquid medium, stationary phase cultures were prepared by inoculating 2.5ml TSB-Kn with frozen S. aureus RN4220 transformants containing phage 77 ORFs cloned in pTHA vector followed by incubation for 16 hrs at 37°C. These cultures were then diluted 1/100 in the same medium, and the bacteria were allowed to grow for 2 hrs at 37°C to reach early log phase. 150 µl of such culture were then mixed with 2.35 ml TSB-Kn medium with or without arsenite (the final concentration of arsenite in the medium was 0 or 5  $\mu$ M arsenite). After 3.5 hrs incubation at 37°C with shaking at 250 rpm, 100 µl of bacterial culture was removed from each tube for OD<sub>565</sub> measurement. Serial ten-fold dilutions of the culture in buffered saline solution (0.85% NaCl) were then spotted onto TSB-Kn plates. The plates were incubated at 37°C 16 hrs and the number of surviving colonies counted the following day. The growth inhibitory property of individual ORFs was then quantitated by comparing CFU numbers under normal or arsenite-induction conditions. A schematic flow of the inhibition analysis is shown in Fig. 3 (also applicable to inhibition analysis for the other phage and bacteria pointed out herein). Inhibition results are shown in Figures 4A-C.

Example 6: Itentification of Cecropin Signature Motif in Staphylococcus aureus

Bacteriophage 3A ORF

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The genome for S. aureus bacteriophage 3A was determined and the sequence was analyzed essentially as described for bacteriophage 77 in the examples above.

Upon blast analysis of the identified open reading frames of phage 3A, the presence of an amino acid sequence corresponding to a cecropin signature motif was observed.

This motif (WDGHKTLEK) is located at position aa 481-489. Cecropins were originally identified in proteins from the cecropia moth and are recognized as potent antibacterial proteins that constitute an important part of the cell-free immunity of insects. Cecropins are small proteins (31-39 amino acid residues) that are active against both Gram-positive and Gram-negative bacteria by disrupting the bacterial membranes. Although the mechanisms by which the cecropons cause cell death are not fully understood, it is generally thought to involve channel formation and membrane destabilization.

The identification of a motif corresponding to a known inhibitor suggests that the product of ORF002 is also an inhibitory compound. Such inhibitory activity can be confirmed as described herein or by other methods known in the art. Confirmation of the inhibitory activity would indicate that the ORF product could serve as the basis for construction of mimetic compounds and other inhibitors directed to the target of the ORF002 product.

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## Example 7. Growth of Staphylococcus aureus bacteriophage 44AHJD:

Staphylococcus aureus propagating strain (PS 44A) (Felix d'Herelle Reference Centre #HER 1101) was used as a host to propagate its respective phage 44AHJD (Felix d'Herelle Reference Centre #HER 101). Two rounds of plaque purification of phage 44AHJD were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the Staphylococcus aureus PS strain was grown overnight at 37°C in Nutrient Broth [NB: 3 g Bacto Beef Extract, 5 g Bactopeptone per liter, (Difco Laboratories # 0003-17-8), supplemented with 0.5% NaCl]. The culture was then diluted 20 fold in NB and incubated at 37°C until an OD<sub>540</sub> of 0.2. In order to obtain single plaques, phage 44AHJD was subjected to 10-fold serial dilutions using the phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin) and 10 μl
were used to infect 0.5 ml of the cell suspension in the presence of 400 μg/ml of

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CaCl<sub>2</sub>. After incubation of 15 min at room temperature, 2 ml of melted soft agar (NB supplemented with 0.6% of agar) were added to the mixture and poured onto the surface of 100 mm nutrient agar plates (3 g Bacto Beef extract, 5 g Bactopeptone, 0.5% NaCl and 15 g of Bacto agar per liter (Difco Laboratories # 0001-17-0). After overnight incubation at 37°C, a single plaque was isolated, resuspended in 1ml of phage buffer by end over end rotation for 2 h at room temperature and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock.

Large scale purification of bacteriophage and preparation of phage DNA was as follows.

The propagation method was carried out by using the agar layer method described by Swanstörm and Adams (1951). Briefly, the PS 44A strain was grown to stationary phase overnight at 37°C in Nutrient Broth. The culture was then diluted 20x in NB and incubated at 37°C until the  $A_{540}$ = 0.2. The suspension (15x10<sup>7</sup> Bacteria) was then mixed with 15x10<sup>5</sup> phage particles to give a ratio of 100-bacteria/phage particle in the presence of 400 µg/ml of CaCl<sub>2</sub>. After incubation of 15 min at room temperature, 7.5 ml of melted soft agar were added to the mixture and poured onto the surface of 150 mm nutrient agar plates and incubated overnight at 37°C. To collect the lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide and shaken vigorously for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) is collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, 10% (w/v) of PEG 8000 and 0.5 M of NaCl were added to the lysate and the mixture was incubated on ice for 16 h. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman).

The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a preformed cesium chloride step gradient as described in Sambrook *et al.* (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 x g) for 24 h at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl<sub>2</sub>. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

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in 20 µl of H<sub>2</sub>O.

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#### Example 8. DNA sequencing of the Bacteriophage 44 AHJD genome.

Four mg of phage DNA was diluted in 200 µl of TE pH 8.0 in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles and size fractionated on 1% agarose gels. The sonicated DNA was then size fractionated by gel electrophoresis. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a coommercial DNA extraction system according to the instructions of the manufacturer (Qiagen) and eluted in 50 µl of 1mMTris-HCl [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymearse and the Klenow fragment of *E. coli* DNA polymerase 1 as follows. Reactions were performed in a final volume of 100 μl containing DNA, 10 mM Tris-HCl pH 8.0, 50 mM NaCl, 10 mM MgCl<sub>2</sub>, 1 mM DTT, 5 μg BSA, 100 μM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of Klenow fragment (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was ethanol precipitated and resuspended

Cloning of the sonicated phage DNA into pKSII vector and transformation:

Blunt-ended DNA fragments were cloned by ligation directly into the *HincII* site of the pkSII vector (Stratagene) dephosphorylated with calf intestinal alkaline phosphatase (New England Biolabs). A typical reaction contained 100 ng of vector, 2

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to 5  $\mu$ l of repaired sonicated phage DNA (50-100 ng) in a final volume of 20  $\mu$ l containing 800 units of T4 DNA ligase (New England Biolabs) overnight at 16°C. Transformation and selection of positive clones was performed in the host strain DH10  $\beta$  of *E. coli* using ampicillin as a selective antibiotic as described in Sambrook et al. (1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 ml LB and 100 µg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *HincII* cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 µl reaction volume containing 10 mM Tris-HCl (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 mM primer, 187.5 µM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep<sup>TM</sup> spin miniprep kit (Qiagen).

using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism BigDye<sup>TM</sup> primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism BigDye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye<sup>TM</sup> terminator cycle sequencing ready reaction kit.

#### Example 9. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher™ 3.1 software

(GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI

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prism BigDye<sup>™</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Staphylococcus aureus* bacteriophage 44AHJD is shown in Table 16.

A software program was used on the assembled sequence of bacteriophage 44AHJD to identify all putative ORFs larger than 33 codons. The software scans the 5 primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG. GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(http://www.ncbi.nlm.nih.gov/htbin-10 post/Taxonomy/wprintgc?mode=c) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, 15 then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage 44AHJD are listed in Tables 17 & 18.

Sequence homology searches for each ORF were carried out using an implementation of blast programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
- ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
- 25 iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
  - iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
  - v) Staphylococcus aureus NCTC 8325 (ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa);
  - vi) Staphylococcuspyogenes (ftp://ftp.tigr.org/pub/data/s\_pneumoniae/gsp.contigs.1121 97.Z);
  - vii)PRODOM(ftp://ftp.toulouse.inra.fr/pub/prodom/current\_release/prodom99\_1.forbl ast.gz);
  - viii) DOMO (ftp://ftp.infobiogen.fr/pub/db/domo/);

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ix) TREMBL (ftp://www.expasy.ch/databases/sp tr nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 44AHJD are shown in Tables 19 & 20.

#### 5 Example 10. Sub-Cloning of Bacteriophage 44 AHJD ORFs.

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 44 AHJD ORF sequence is inducible. For example, the shuttle vector pT0021, in which the firefly luciferase (lucFF) expression is controlled by the ars (arsenite) promoter/operator (Tauriainen et al., 1997), can be modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the heamaglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with BamHI, SalI and HindIII cloning sites) is: 5'-gatcccggtcgaccaagcttTACCCATACGACGTCCCAGACTACGCCAGCTGA-3' (where upper case letters denote the nucletotide sequence of the HA tag); the antisense strand HA tag sequence (with a HindIII cloning site) is: 5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAaagcttggtcgaccgg-3' (where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with BamHI and HindIII. This manipulation resulted in replacement of the lucFF gene by the HA tag. This modified shuttle vector containing the arsenite inducible promoter, the arsR gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A (another userful vector construct is shown in Fig. 1B).

Each ORF, encoded by Bacteriophage 44 AHJD, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon can be selected for functional analysis for bacterial inhibition. Each individual ORF, from initiation codon to last codon (excluding the stop codon), can be amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of ORFs, each sense strand primer targets the initiation codon and is preceded by a BamHI restriction site (scggatcc3) and each antisense oligonucleotide targets the pentultimate codon (the one before the stop codon) of the ORF and is preceded by a Sal I restriction site (scggatcgaccg3). The PCR product of each ORF can be gel

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purified and digested with BamHI and SaII. The digested PCR product can then be gel purified using the Qiagen kit as described, ligated into BamHI and SaII digested pTHA vector, and used to transform E. coli bacterial strain DH10β(as described above). As a result of this manipulation, the HA tag is set inframe with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones will be picked and their insert sizes were confirmed by PCR analysis using primers flanking the cloning site. The following primers can be used for PCR amplification: HAF: 5'TATTATCCAAAACTTGAACA<sup>3'</sup>; HAR: 5'CGGTGGTATATCCAGTGATT<sup>3'</sup>. The sequence integrity of cloned ORFs can be verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence can not be achieved by one pass with the sequencing primers, additional internal primers will be selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiswirth et al., 1983) will be used as a recipient for the expression of recombinant plasmids. Electoporation will be performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones will be performed on Luria-Broth agar (LB-agar) plates containing 30 µg/ml of kanamycin.

Alternatively, a constitutive promoter can be used to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids will be introduced into *Staphylococcus aureus* strain RN4220 (Kreiswirth et al., 1983) using electoporation as previously described (Schenk and Laddaga, 1992).

#### Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), can be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10. Recombinant clones are then picked and their insert sizes confirmed by

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PCR analysis using primers flanking the cloning site as well as restriction digestion. The sequence fidelity of cloned ORFs can be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal primers can be selected and used for sequencing. Recombinant plasmids can be introduced into Staphylococcus aureus strain RN4220 (Kreiswirth et al., 1983) using electoporation as previously described (Schenk and Laddaga, 1992).

Induction of gene expression from the ars promoter.

If an inducible promoter is used, e.g., the ars promoter, induction can be assessed, for example, in either of the two methods.

#### 1. Screening on agar plates

The functional identification of killer ORFs can be performed by spreading an aliquot of S. aureus transformed cells containing phage 44 AHJD ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 µM). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite. 2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase (OD<sub>540</sub>=.2) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 µl/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 µM) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage 44 AHJD ORFs on bacterial cell growth is then monitored by measuring the OD<sub>540</sub> and comparing the rate of growth to the culture not containing inducer. [As positive controls for growth inhibition, the kilA gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 Virology #193: 1033-1036), and the holin/lysin genes of the Sthaphylococcus aureus phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G., Maier, SK. and Scherer, S. 1998. FEMS Microbiology Letters #162:265-274) can be subcloned into the ars inducible vector. An aliquot of the induced and uninduced... culture can also be plated out on agar plates containing an appropriate antibioticselection but lacking inducer. Following incubation overnight at 37°C, the number of

colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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Example 11. Growth of Enterococcus bacteriophage 182 and purification of genomic DNA.

The Enterococcus propagating strain (PS) (Enterococcus sp. Group D, Felix d'Herelle Reference Centre #HER 1080) was used as host to propagate its respective phage 182 (Felix d'Herelle Reference Centre #HER 80). Two rounds of plaque purification of phage 182 were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the Enterococcus sp. PS strain was grown overnight at 37°C in Tryptic Soy Broth [TSB: 17 g Bacto tryptone, 3 g Bacto soytone, 2.5 g Bacto dextrose, 5 g Sodium chloride, and 2.5 g Dipotassium phosphate per liter (Difco Laboratories (#0370-17-3)]. The culture was then diluted 20 fold in TSB and incubated at 37°C until the OD<sub>540</sub>= 0.2 (early log phase) with constant agitation. In order to obtain single plaques, phage 182 was subjected to 10 fold serial dilutions using the phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 l of each dilution was used to infect 0.5 ml of the bacterial cell suspension. After incubation at 15 min at 37°C, 2 ml of melted soft agar (TSB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm Trytic Soy Agar plates [TSA: 15 g Tryptone peptone, 5 g Soytone peptone, 5 g Sodium chloride and 15 g of Agar per liter (Difco Laboratories #0369-17)]. After overnight incubation at 37°C, a single plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

The propagation procedure for bacteriophage 182 was modified from the agar layer method of Swanstörm and Adams (1951). Briefly, the *Enterococcus* sp. PS strain was grown to stationary phase overnight at 37°C in TSB. The culture was then diluted 20 fold in TSB and incubated at 37°C until the  $A_{540}$ = 0.2. The suspension (15x10<sup>7</sup> Bacteria) was then mixed with 15x10<sup>5</sup> plaque forming units (pfu) to give a

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ratio of 100-bacteria/pfu. After incubation of 15 min at 37°C, 7.5 ml of melted soft agar (TSB plus 0.6% agar) were added to the mixture and poured onto the surface of 150 mm TSA plates and incubated 16 hrs at 37°C. To collect the plate lysate, 20 ml of TSB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant fluid (lysate) is collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) of PEG 8000 and 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phages were harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl<sub>2</sub>. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 g/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

### Example 12. DNA sequencing of the Bacteriophage 182 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4

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cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 μl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl<sub>2</sub>, 1 mM DTT, 50 μg/ml BSA, 100 μM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of the Klenow large fragment of DNA polymerase I(New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in 20 μl of H<sub>2</sub>O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook *et al.*, 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μl LB and 100 μg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the Hinc II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 μl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 μM primer, 187.5 μM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec

denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the OIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye<sup>TM</sup> primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye<sup>TM</sup> terminator cycle sequencing ready reaction kit.

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### Example 13. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher<sup>™</sup> 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI prism BigDye<sup>™</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Enterococcus* bacteriophage 182 is shown in Table 21.

A software program was used on the assembled sequence of bacteriophage 182 to identify all putative ORFs larger than 33 codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(http://www.ncbi.nlm.nih.gov/htbin-

30 <u>post/Taxonomy/wprintgc?mode=c</u>) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the

next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is

- performed on all three reading frames of both DNA strands of the phage sequence.

  The predicted ORFs for bacteriophage 182 are listed in Tables 22 & 23.

  Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:
- 10 (i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
  - ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
  - iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
  - iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
  - v) staphylococcus aureus NCTC 8325 (ftp://ftp.genome.ou.edu/pub/staph/staph-
- 15 lk.fa);

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- vi) streptococcus pyrogenes
- (ftp://ftp.tigr.org/pub/data/s pneumoniae/gsp.contigs.112197.Z);
- vii) PRODOM
- (ftp://ftp.toulouse.inra.fr/pub/prodom/current\_release/prodom99.1.forblast.gz);
- viii) DOMO (ftp://ftp.infobiogen.fr/pub/db/domo/);
  - ix) TREMBL (ftp://www.expasy.ch/databases/sp\_tr\_nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 182 are shown in Tables 24 & 26.

#### 25 Example 14. Sub-Cloning of Bacteriophage 182 ORFs.

#### Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 182 ORF sequence is inducible. For example, the plasmid pND50 replicates in *E. coli*, *E. faecalis*, and *S. aureus* (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. *Antimocrob. Agents Chemother*. 40, 1157-1163). This plasmid—can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the firefly luciferase (*lucFF*)

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expression is controlled by the *ars* promoter/operator from a *S. aureus* plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997). Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. *Appl. Environ. Microbiol.* 63:4456-4461). This modified shuttle vector will contain the *ars* promoter, *arsR* gene and a cloning site for introduction of individual phage ORFs downstream from a shine-delgarno sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system The nisA promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the nisA promoter (10- to 60-fold induction) can be obtained in all of the species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transciption in *Enterococcus*.

Alternatively, a constitutive promoter can be used (e.g., the β-lactamase promoter is constitutive in *E. faecalis* – see ref. 1) to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *E. faecalis* strain FA2-2 by electroporation, as previously described (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. *Antimicrob. Agents Chemother*. 40, 1157-1163). Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgamo sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), will be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on

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the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10 $\beta$ . Recombinant clones are then picked and their insert sizes confirmed by PCR analysis using primers flanking the cloning site as well as restriction digestion. The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into E. faecalis strain FA2-2 by electroporation, as previously described

(Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. Antimicrob. Agents Chemother. 40, 1157-1163).

Induction of gene expression from the ars promoter.

If an inducible promoter is used, e.g., the ars promoter, induction can be assessed, for example, in either of the two methods.

### 15 <u>1. Screening on agar plates</u>

The functional identification of killer ORFs can be performed by spreading an aliquot of *E. faecalis* transformed cells containing phage 182 ORF onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 µM). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite.

#### 2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase (OD<sub>540</sub>=.2) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 μl/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μM) and the culture incubated for an additional 2 h at 37°C. The effect of expression of the phage 182 ORFs on bacterial cell growth is then monitored by measuring the OD<sub>540</sub> and comparing the rate of growth to the culture not containing inducer. As positive controls for growth inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lysin* genes of the *Sthaphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G.,

Maier, SK. and Scherer, S. 1998. FEMS Microbiology Letters #162:265-274) were subcloned into the ars inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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# Example 15. Growth of Streptococcus bacteriophage Dp-1 and purification of genomic DNA.

The Streptococcus pneumoniae R6 propagating strain (PS) (Tomasz, 1966) was used as host to propagate its respective phage Dp-1 (McDonnell et al., 1975). (Alternatively, Streptococcus (Diplococcus) pneumoniae R36A could be used. Strain R36A is available from ATCC as #11733 or 27336. Streptococcus pneumoniae is also available from Felix d'Herelle Reference Center in Quebec, Canada as catalog number HER 1054. Other S. pneumoniae strains are also available from ATCC.)

Two rounds of plaque purification of phage Dp-1 were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the Streptococcus R6 PS strain was grown overnight at 37°C in K-Cat media [K-Cat: 10 g Bacto casitone, 5 g Bacto tryptone, 1 g Yeast extract, 5g Potassium chloride, 0.2% Glucose, 30mM Potassium phosphate buffer [pH 8] and 250,000 Units Catalase per liter (Boehringer Mannheim #10683600). The culture was then diluted 20 fold in K-CAT and

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incubated at 37°C until the OD<sub>540</sub>= 0.2 (early log phase) with constant agitation. In order to obtain single plaques, Dp-1 phage was subjected to 10-fold serial dilutions using the phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM MgCl<sub>2</sub>)and 10 μl of each dilution was used to infect 0.5 ml of the cell suspension.
5 After incubation of 15 min at 37°C, 2 ml of melted soft agar (K-CAT supplemented with 0.8% of agar) were added to the mixture and poured onto the surface of 100 mm K-CAT agar plates [K-CAT supplemented with 1.2 % of agar]. After solidification of the soft agar layer, an additional 5 ml of melted soft agar was added to visualize distinct plaques (Ronda et al., 1978). After overnight incubation at 37°C, a single plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

The propagation procedure for bacteriophage Dp-1 was modified from the agar layer method of Swanstörm and Adams (1951). Briefly, the R6 strain of Streptococcus pneumoniae was grown to stationary phase overnight at 37°C in K-CAT. The culture was then diluted 20 fold in K-CAT and incubated at 37°C until the  $OD_{540} = 0.2$ . The suspension  $(15x10^7 \text{ Bacteria})$  was then mixed with  $15x10^5$  plaque forming units (pfu) to give a ratio of 100-bacteria/pfu. After incubation of 15 min at 37°C, 7.5 ml of melted soft agar (K-CAT plus 0.8% agar) were added to the mixture and poured onto the surface of 150 mm K-CAT agar plates and incubated 16 hrs at 37°C. After solidification of the soft agar layer, 7.5 ml of melted soft agar were added to each plate. To collect the plate lysate, 20 ml of K-CAT media were added to each plate and the soft agar layers were collected by scrapping off with a clean microscope slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) was collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) of PEG 8000 and 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM MgCl<sub>2</sub>). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS-55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl<sub>2</sub>. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

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#### Example 16. DNA sequencing of the Bacteriophage Dp-1 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 sec spaced by 15 sec cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume,  $100~\mu l$ ) containing sonicated phage DNA, 10~mM Tris-HCl [pH 8.0], 50~mM NaCl, 10~mM MgCl<sub>2</sub>, 1~mM DTT,  $50~\mu g/ml$  BSA,  $100~\mu M$  of each dNTP and 15~units of T4 DNA polymerase (New England Biolabs) for 20~min at  $12^{\circ}$ C followed by addition of 12.5~units of the Klenow large fragment of DNA polymerase I (New England Biolabs) for 15~min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in  $20~\mu l$  of  $H_2O$ .

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection

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of bacterial clones containing recombinant plasmids was performed in E. coli DH10β according to standard procedures (Sambrook et al., 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μl LB and 100 μg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 μl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 μM primer, 187.5 μM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep<sup>TM</sup> spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye<sup>TM</sup> primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit.

Example 17. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher<sup>TM</sup> 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI prism BigDye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Streptococcus* bacteriophage Dp-1 is shown in Table 28.

A software program was used on the assembled sequence of bacteriophage

Dp-1 to identify all putative ORFs larger than 33 codons. The software scans the

primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon.

Three possible selections can be made for defining the nature of the start codon; I)

selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG,

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GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(<a href="http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c">http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c</a>) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage Dp-1 are listed in Tables 29 and 30, and Fig. 6.

Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
- ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
- iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
- iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
- v) staphylococcus aureus NCTC 8325
- 20 (ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa);
  - vi) streptococcus pyogenes

(ftp://ftp.tigr.org/pub/data/s\_pneumoniae/gsp.contigs.112197.Z);

vii) PRODOM

(ftp://ftp.toulouse.inra.fr/pub/prodom/current\_release/prodom99.1.forblast.gz);

- viii) DOMO (ftp://ftp.infobiogen.fr/pub/db/domo/);
- ix) TREMBL (ftp://www.expasy.ch/databases/sp\_tr\_nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage Dp-1 are shown in Table 31.

Example 18. Sub-Cloning of Bacteriophage Dp-1 ORFs.

#### Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage Dp-1 ORF sequence is inducible.

For example, the plasmid pLSE4 replicates in *E. coli*, and *S. pneumoniae* (Diaz and Garcia, 1990). This plasmid can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the

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firefly luciferase (lucFF) expression is controlled by the ars promoter/operator from a S. aureus plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997). Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. Appl. Environ. Microbiol. 63:4456-4461). This modified shuttle vector will contain the ars promoter, arsR gene and a cloning site for introduction of individual phage ORFs downstream from a shine-dalgamo sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system The nisA promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the nisA promoter (10- to 60-fold induction) can be obtained in all of the species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transcription in *Streptococcus*.

Alternatively, a constitutive promoter can be used to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *S. pneumoniae* R6 as previously described (Diaz and Garcia, 1990)

### Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), will be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10\(\textit{B}\). Recombinant clones are then picked and their insert sizes confirmed by PCR analysis using primers flanking the cloning site as well as restriction—digestion. The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site

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internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into S. pneumoniae R6 as previously described (Diaz and Garcia, 1990). Induction of gene expression from the ars promoter.

If an inducible promoter is used, e.g., the ars promoter, induction can be assessed, for example, in either of the two methods.

## 1. Screening on agar plates

The functional identification of killer ORFs can be performed by spreading an aliquot of S. pneumoniae transformed cells containing phage Dp-1 ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5  $\mu$ M). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite.

## 2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are 15 then diluted to the mid log phase (OD<sub>540</sub>=.2) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 µl/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 µM) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage Dp-1 ORFs on bacterial cell growth is then monitored by measuring the OD<sub>540</sub> and comparing the rate 20 of growth to the culture not containing inducer. [As positive controls for growth inhibition, the kilA gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 Virology #193: 1033-1036), and the holin/lysin genes of the Sthaphylococcus aureus phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G., Maier, SK. and Scherer, S. 1998. FEMS Microbiology Letters #162:265-274) can be 25 subcloned into the ars inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of 30 inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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All patents and publications mentioned in the specification are indicative of the levels of skill of those skilled in the art to which the invention pertains. All references cited in this disclosure are incorporated by reference to the same extent as if each reference had been incorporated by reference in its entirety individually.

One skilled in the art would readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. The specific methods and compositions described herein as presently representative of preferred embodiments are exemplary and are not intended as limitations on the scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention are defined by the scope of the claims.

It will be readily apparent to one skilled in the art that varying substitutions and modifications may be made to the invention disclosed herein without departing from the scope and spirit of the invention. For example, those skilled in the art will recognize that the invention may suitably be practiced using a variety of different bacteria, bacteriophage, and sequencing methods within the general descriptions provided.

The invention illustratively described herein suitably may be practiced in the absence of any element or elements, limitation or limitations which is not specifically disclosed herein. Thus, for example, in each instance herein any of the terms "comprising," "consisting essentially of and "consisting of may be replaced with either of the other two terms. The terms and expressions which have been employed are used as terms of description and not of limitation, and there is not intention that in the use of such terms and expressions of excluding any equivalents of the features shown and described or portions thereof, but it is recognized that various modifications are possible within the scope of the invention claimed. Thus, it should be understood that although the present invention has been specifically disclosed by preferred embodiments and optional features, modification and variation of the concepts herein disclosed may be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of this invention as defined by the appended claims.

In addition, where features or aspects of the invention are described in terms of Markush groups or other grouping of alternatives, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group or other group. For example, if there are alternatives A, B, and C, all of the following possibilities are included: A separately, B separately, C separately, A and B, A and C, B and C, and A and B and C. Thus, for example, for the bacteria and phage specified herein, the embodiments expressly include any subset or subgroup of those bacteria and/or phage. While each such subset or subgroup could be listed separately, for the sake of brevity, such a listing is replaced by the present description.

Thus, additional embodiments are within the scope of the invention and within the following claims.

Table 1

# Phages against human and animal pathogenic bacteria

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I. Pathogen name	Phage name	II.	Cat alo g#	Origin/reference
Acinetobacter calcoaceticus	A3/2 A10/45 A36 B9GP B <sub>9</sub> PP BS46 E13		<u>gr</u>	Felix d'Herelle Reference Centre,Quebec,Quebec
	E14 531 Ap3			J. Bacteriol 1984. 157: 179-183
Acinetobacter haemolyticus	P78			J. Gen. Microbiol 1986.132: 2633-2636  Felix d'Herelle Reference Centre, Quebec, Quebec
Acinetobacter johnsonii				Felix d'Herelle Reference Centre.Quebec,Quebec
Acinetobacter sp.	BP1 G4, HP2, HP3 & HP4			J.Virol.1968.2:716-722  Can.J.Microbiol.1966.12:1023-1030 & J.Virol.1974.13:46-52 & Arch.Virol.1994.135:345-354
	A1, A4, A9 & 196			Arch. Virol. 1994.135:345-354
	HP1 A19, A23, A29, A31, A33, A34, A3759 & 2845			Can.J.Microbiol.1966.12:1023-1030  J.Microsc (Paris) 1973.16:215-224 &  CR.Hebdo Seances Acad.Sci.Ser D.Sci Natur(Paris)278:1907-1909 &  Arch.Virol.1994.135:345-354 &  Rev.Can.Biol.1970.29:317-320
Actinobacillus actinomycetecomitans				FEMS Microbiol Lett 1994. 119:329-337

			Infec. Immun. 1982. 35: 343-349
		<b>[</b>	
			Mol.Gen.Genet 1998.258: 323-325
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		+	0.136
	Ααφ247	10000	Oral Micriol. Immunol 1997.12: 40-46
Actinomyces viscosus	İ	43146-B1	The American Type Culture Collection
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			Infect.Immun.1985.48:228-233
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			Infect.Immun.1988.56:54-59
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		1	Plasmid 1997.37:141-153
		1	
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Aeromonas hydrophila	PM2** & PM3		FEMS Microbiol.Lett. 1990.57:277-282
	Aehl	1	Felix d'Herelle Reference
	Aeh2		Centre,Quebec,Quebec
	PM4		<u> </u>
	PM5		
	РМ6		
	T7-ah		
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Aeromonas	3	T	Felix d'Herelle Reference
salmonicida	25		Centre, Quebec, Quebec
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	Asp37		Com I Missabial 1092 20: 1459 1461
	55R.1	07005 D1	Can. J. Microbiol. 1983. 29: 1458-1461
Alteromonas espejiana	PM2**	27025-B1	The American Type Culture Collection
Asticacaulis			Felix d'Herelle Reference
biprosthecum			Centre, Quebec, Quebec
Asticcacaulis		15261-B1	The American Type Culture Collection
excentricus		15261-B2	
		15261-B3	
		1020. 25	
	фАс21	1	
	фАс24		
Azotobacter vinelandii		12518-B1	The American Type Culture Collection
		12518-B4	
		12518-B5	·
	A14	12518-B9	
	A21	12518-B10	
	A31	13705-B1	
	A41		
	PAV1		711 1 1050 40 420 450
Azotobacter sp.			Virology 1972.49:439-452
Bacteroides fragilis	Bf-1		Rev. Infect. Dis. 1979. 1: 325-336
· · · · · · · · · · · · · · · · · · ·	B40-8		FEMS Microbiol. Lett. 1991. 66: 61-67
	HSP40		Appl. Environ. Microbiol. 1989. 55: 2696- 2701
	phiA1		Zentralbl.bakteriol.1972.222:57-63
Bdellovibrio	MAC-1		J. Gen. Microbiol. 1987. 133: 3065-3070
bacteriovorus			
Bdellovibrio sp.	VL-1		J.Virol.1973.12:1522-1533
Bordetella brochiseptica	214		Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-

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Bordetella			Felix d'Herelle Reference
parapertussis			Centre, Quebec, Quebec
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			Mol. Gen. Mikrobiol. Virusol. 1988.4: 22-25
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			Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-
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	41403		13
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Brucella abortus	<b>'</b>		Centre, Quebec, Quebec
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			- Calleria
		23448-B1	The American Type Culture Collection
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		23448-B3	
	}	17385-B1	
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	DIZ 2 TD 0		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2:
	BK-2, TB &		48-52
	Fi		
	R/c & R/O	ļ	Dev. Biol. Stand. 1984.56: 55-62
Brucella canis	R/c		Dev. Biol. Stand. 1984.56: 55-62
Brucella melitensis	BK-2	23456-B1	The American Type Culture Collection
Brucella suis	Wb		Zentralbl. Veterinarmed. 1975.22:866-867
Drucena suis	""		
	1	1	1

	Fi** & TB		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2: 48-52
Brucella sp.			Can. J. Vet. Res. 1989.53: 319-325
			Res. Vet. Sci. 1988. 44: 45-49
			Res. Vel. Bel. 1900. VI. 15
	R		Zh.Mikrtobiol.Epidemiol.Immunobiol.1983.2:
Campylobacter coli		43133-B1	The American Type Culture Collection
	10	43134-B1	The American Time Cultura Collection
Campylobacter coli	18	43135-B1	The American Type Culture Collection
(Cont'd)	19 20	43136-B1	
Campylobacter jejuni	1	35918-B1	The American Type Culture Collection
Campyioodeioi jojami	2	35919-B1	
	3	35920-B1	
	4	35921-B1	
	5	35918-B2	
	6	35920-B2	
	7	35922-B2	ŀ
	į.	35923-B1	1
	8	35924-B1	
	9	35925-B1	
	10	35925-B2	
	11	35922-B2	
	12	35924-B2	1
	13		
	14	35922-B3	
	17	43133-B1	
	18	43134-B1	
	19	43135-B1	
	20	43136-B1	
Campylobacter (Helicobacter) pylori	HP1		J. Med. Microbiol.1993. 38: 245-249
Chlamydia psittaci	Chp1**		J. Gen. Virol. 1989. 70: 3381-3390
Clostridium	CAK-1		J.Bacteriol.1993.175:3838-3843
acetobutylicum			

Clostridium botulinum	T	Nucleic Acids Res.1990.18:1291
		Bioch.Biophys.res.Commun.1990.171.1304- 1311
		Microbiol.immunol.1981.25:915-927
		J.Vet.Med.Sci.1992.54:675-684
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	СЕ β & СЕ γ	
Clostridium difficile	41 & 56	J. Clini.Microbiol. 1985.21:251-254

Clostridium Rev.Can.Biol.1977.36:205-2	15
FEMS Microbiol.Lett. 1990.	54:323-326
	Collection
Clostridium         8074-B1         The American Type Culture of 17886-B1           59         17886-B1         70           17886-B3         71         17886-B4           72S         17886-B5         72L           17886-B6         17886-B6	Concenton
Clostridium tetani A & B Rev.Can.Biol.1978.37:43-46	
Corynebacterium diphteriae Vopr.Virusol.1986.31:577-58	
Corynebacterium NN 12319-B1 The American Type Culture	Collection
pseudotuberculosis	Collection

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Enterococcus faecalis	42	19948-B1	The American Type Culture Collection
Enterococcus faecium		19950-B1	The American Type Culture Collection
		19953-b2	
		19953-B1	
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Escherichia coli		11303-B14	The American Type Culture Collection
		11303-B10	
		11303-B21	
		8677-B1	
		11303-B13	
	}	13706-B4	
Escherichia coli		15766-B1	The American Type Culture Collection
(Cont'd)	İ	15766-B1	
		1242-B5	
		15669-B2	
		15767-B1	
		11303-B16	
		27-65-B1	
	C204	25065-B2	
	E1	15669-B1	
	fi**	15597-B1	
	f2**	21816-B1	
	l l	23724-B9	
	FCZ fd**	15593-B1	:
	Id**	25404-B1	
		29746-B1	
		23631-B1	
		25868-B1	
		25298-B1	
		25298-B2	
		11303-B37	
		11303-B24	
	IU**	11303-B26	
	111	11303-B27	
		11303-B28	
		11303-B29	
		11303-B30	
		11303-B33	
		11303-B31	
		11303-B25	
		11303-B35	
	MS2**	11303-B34	
	MU9	11303-B36	
	Mu-1	11303-B32	
	Ox6	13706-B5	
	P1**	11303-B1	
	P4 sid; **	11303-B2	
	Q-β**	11303-B3	
	R17**	11303-B4	
	Z1K/1	35060-B1	
	ZJ/2	35060-B2	
	2212	35060-B3	-
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		11303-B7	
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Escherichia coli		11303-B20	The American Type Culture Collection
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		11303-B15	·
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	547	11303-B18	
	UVI	13706-B2	
	UV47	23724-B2	
	UV375	23724-B1	
	α3 <sup>‡‡</sup>	23724-B3	
	λ **	23724-B4	
	λ C-17	23724-B5	
	λ sus P-3	23724-B6	
	λ sus R-5	23724-B7	
	λ sus J-6	23724-B8	· ·
	λ sus O-8	35860-B1	
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	λ ind	15597-B2	
	ø92	13706-B1	
	ØR.	49696-B1	
	øV-1	Ì	}
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	K18 & Oxl M1°°, TuIa°° & TuIb°°		J.Mol.Biol.1987.196:165-174
	K10		J.Bacteriol.1979.140:680-686
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	PRD1		
	K3hx		Mol.Gen.Genet.1987.206:110-115
	933J**& 933W**		Infect.Immunity.1986.53:135-140
	H19-B**		J.Bacteriol.1987.169:4308-4312
	Tcp-111		Zentralbnl.Bakteriol.Mikrobiol.Hyg.1988.27 41-51

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Obeta 1		N4**	Vet.Microbiol.1992.30:203-212
Obeta 1   J.Bacteriol.1978.133:172-177     P1CM		Phi 80 trp	Ann.Inst.Pasteur.1971.120:121-125
PA-2**   J.Bacteriol.1990.172:1660-1662     186**   Mol.Gen.Genet.1982.187:87-95     186.IX.B   Mol.Microbiol.1992.6:2629-2642     21**   Virology 1983.129:484-489     P4**   MicrobiolRev.1993.57:683-702     82**   J.Biol.Chem.1987.262:11721-11725     PSP3   J.Bacteriol.1996.178:5668-5675     HK022**   Nucleic Acids Res.1994.22:354-356     D108**   Nucleic Acids Res.1986.14:3813-3825     Escherichia coli (Cont'd)   Re**   J.Mol.Biol.1997.267:237-249     (Cont'd)   Re**   J.Mol.Biol.1985.181:27-39     P22dis   Mol.Gen.Genet.1978.166:233-243     N15**   J.Bacteriol.1996.178:1484-1486     If1**   Proc.R.Soc.Lond.B.Biol.Sci.1991.245:23-30     Stx2Phi-I & Stx2Phi-II     18   Virology 1987.156:122-126     X   J.Gen.Microbiol.1981.126:389-396		Obeta 1	J.Bacteriol.1978.133:172-177
186°*   Mol.Gen.Genet.1982.187:87-95     186.IX.B		PICM	J.Gen.Microbiol.1978.107:73-83
186°°   Mol.Gen.Genet.1982.187:87-95     186.IX.B		PA-2**	J.Bacteriol.1990.172:1660-1662
21**   Virology 1983.129:484-489     P4**   MicrobiolRev.1993.57:683-702     82**   J.Biol.Chem.1987.262:11721-11725     PSP3   J.Bacteriol.1996.178:5668-5675     HK022**   Nucleic Acids Res.1994.22:354-356     D108**   Nucleic Acids Res.1986.14:3813-3825     Escherichia coli (Cont'd)   Re**   J.Mol.Biol.1997.267:237-249     P22dis   Mol.Gen.Genet.1978.166:233-243     N15**   J.Bacteriol.1996.178:1484-1486     If1**   Proc.R.Soc.Lond.B.Biol.Sci.1991.245:23-30     Stx2Phi-I & Stx2Phi-II     18   Virology 1987.156:122-126     X   J.Gen.Microbiol.1981.126:389-396		186**	Mol.Gen.Genet.1982.187:87-95
P4**   MicrobiolRev.1993.57:683-702   82**   J.Biol.Chem.1987.262:11721-11725   PSP3   J.Bacteriol.1996.178:5668-5675   HK022**   Nucleic Acids Res.1994.22:354-356   D108***   Nucleic Acids Res.1986.14:3813-3825   Escherichia coli (Cont'd)   Rb49   J.Mol.Biol.1997.267:237-249   Ike**   J.Mol.Biol.1985.181:27-39   P22dis   Mol.Gen.Genet.1978.166:233-243   N15**   J.Bacteriol.1996.178:1484-1486   If1**   Proc.R.Soc.Lond.B.Biol.Sci.1991.245:23-30   Stx2Phi-II & Stx2Phi-II   Virology 1987.156:122-126   X   J.Gen.Microbiol.1981.126:389-396		186.IX.B	Mol.Microbiol.1992.6:2629-2642
S2**   J.Biol.Chem.1987.262:11721-11725		21**	Virology 1983.129:484-489
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HK022**   Nucleic Acids Res.1994.22:354-356     D108**		82**	J.Biol.Chem.1987.262:11721-11725
HK022**   Nucleic Acids Res.1994.22:354-356     D108**		PSP3	J.Bacteriol.1996.178:5668-5675
D108**   Nucleic Acids Res.1986.14:3813-3825	·		Nucleic Acids Res.1994.22:354-356
Escherichia coli         Rb49         J.Mol.Biol.1997.267:237-249           (Cont'd)         Ike***         J.Mol.Biol.1985.181:27-39           P22dis         Mol.Gen.Genet.1978.166:233-243           N15**         J.Bacteriol.1996.178:1484-1486           If1**         Proc.R.Soc.Lond.B.Biol.Sci.1991.245:23-30           Stx2Phi-I         Mol.Gen.Genet.1978.166:233-243           Virology 1987.156:122-126         Virology 1987.156:122-126           X         J.Gen.Microbiol.1981.126:389-396			Nucleic Acids Res.1986.14:3813-3825
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	TYDIAA		Nucleic Acids Res. 1996.24:2360-2368
Haemophilus influenzae	HP1**		Gene 1997. 196: 139-144
	S2**		Felix d'Herelle Reference
Halobacterium cutirubrum	S45		Centre, Quebec, Quebec
Halobacterium			Felix d'Herelle Reference
halobium			Centre, Quebec, Quebec
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			Can.J.Microbiol.1982.28:916-921
	1		Qui.3.1711V2QQ1Q1.17Q2.20171Q-721
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Halobacterium			Biol.Chem.Hoppe Seyler 1994.375:747-757
salinarium	1	1	
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Klebsiella oxytoca	tf-1		J.Gen.Microbiol.1987.133:953-960
Klebsiella pneumoniae	60 92	23356-B1 23357-B1	The American Type Culture Collection
	K19Q		Felix d'Herelle Reference
			Centre, Quebec, Quebec
	FC3-1 & FC3-9		Can.J.Microbiol.1991.37:270-275
	FC3-10		FEMS Microbiol.Lett.1991.67:291-297
Klebsiella sp.	K11**		Mol.Gen.Genet. 1990.221:283-286
Leptospira sp.	LE1, LE3 & LE4		Res.Microbiol.1990.141:1131-1138
Listeria	243	23074-B1	The American Type Culture Collection
monocytogenes	197,1313 & 9425		Appl.Environ.Microbiol.1997.63:3374-3377
	H387 & H387-A		Appl.Environ.Microbiol.1993.59:2914-2917
	5775,6223 &12682		APMIS.1993.101:160-167
	2389, 2671,		Intervirology 1994.37:31-35 &
	4211 & 2685	l	Zentralbl.Bakteriol.Mikrobiol.Hyg.1986.261: 2-28
	4b, 4ab, 4g & 3c		Ann.Microbiol (Paris) 1977.128:185-198
	A118, A500 & A511**		Mol.Microbiol. 1995.16:1231-1241-992
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	1/2a, 1/2b, 3c, 4ab, 6a & 6b		Clin.Invest.Med.1984.7:229-232
	φLMUP35 2685		Felix d'Herelle Reference Centre, Quebec, Quebec
Listeria innocua	4211		Felix d'Herelle Reference Centre, Quebec, Quebec
Micrococcus luteus		4698-B1 4698-B4	The American Type Culture Collection
	N3	4698-2	
	N4	4698-B3	
	N8		
Micrococcus luteus	N17		Can.J.Microbiol. 1979.25:1027-1035
Mycobacterium	BK-3	27203-B1	The American Type Culture Collection
smegmatis	Bo1**	27204-B1	
_	Bo 6	27205-B1	
	Bo 6II	27205-B2	
	Bo 6III	27205-B3	
	Mc-2	607-B6	
	Mc-4	607-B7	
	NN	11727-B1	
	Phagus lacticola R1	11759-B1 607-B1	

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	Legendre Leo Roy Sedge		Mol.Microbiol. 1993.7:395-405
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Mycobacterium fortuitum	Bo 4 Bo 7	23052-B1 27207-B1 27207-B2	The American Type Culture Collection

Mycobacterium leprae			Ann.Microbiol. (Paris) 1982.133:93-97
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Mycobacterium		25618-B1	The American Type Culture Collection
tuberculosis	DS6A	25618-B2 4243-B1	
	DSUA		
	110, 139 & 33D		Arch.Virol.1993.133:39-49
	AG1,GS4E,		The Biology of Mycobacteria. Academic
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	PH & BK1	11760 51	1982.309-351
Mycobacterium sp	Phagus pellegrini	11760-B1 11761-B1	The American Type Collection Culture
	NN B1	23239-B1	

Mycobacterium vaccae Mycobacterium phlei	TM4, ph60, ph72, PhAE39, phAE40 & Bxb1 C2 18 & I15 63 phlei & butyricum MyF3P-59a Bo2a D4,D28 & D32 HC B5	15483-B1 11728-B1 11758-B1 27086-B2 27086-B1	Experentia 1969.25:1112-1113 J.Gen.Virol.1987.68:949-956 Gruzlica 1968.36:617-622 J.Gen.Virol.1975.29:235-238  Z.Allg.Mikrobiol.1968.8:29-37 J.Gen.Virol.1973.20:75-87 J.Exptl.Med.1966.123:327-340 J.Bacteriol.1963.86:608-609 The American Type Culture Collection  The American Type Culture Collection
Mycoplasma arthritidis	MAV1**		Infect.Immunity.1995.63:4016-4023
Mycoplasma hyorhinis	Hr-1	<del>                                     </del>	Arch.Virol.1983.77:81-85
Mycoplasma pneumoniae	Br-1		Arch.Virol.1983.75:1-15
Mycoplasma pulmonis			Plasmid 1995. 33: 41-49
Mycoplasma sp.			J.Gen.Microbiol.1985:131:3117-3126
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MV-L2 &	·	Arch.Virol.1979.61:289-296
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Neisseria perflava			J.C.IIIWICIODIOI.1770. 4.07 71
			J.Gen.Virol.1974.23:247-254
Nocardia erythrypolis	φС		J.Bacteriol.1976.126:1104-1107
	φЕС	<del> </del>	Arch.Exp.Veterinarmed.1981.35:433-436
Pasteurella multocida	B225	<u> </u>	Am.J.Vet.Res.1978.39:1565-1566
	B939a		Vet.Med.Nauki. 1977.14:33-36
	Nos.115, 32, 967		V CLIVICU.INDURI. 17/7.14.33-30
	1075		
Propionibacterium	NN	29399-B1	The American Type Collection Culture
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Pseudomonas		12175-B1	The American Type Culture Collection
aeruginosa	2	12175-B2	
	2A	12175-B3	
	2B	12175-B4	
	11	14205-B1	
	16	14206-B1	
	24	14207-B1	
	27	14208-B1	
	44	14209-B1	
	73	14210-B1	
	95	14211-B1	
	109	14212-B1	
	113	14213-B1	
	249	14214-B1	
	B3	15692-B1	
	Hoff 2	14203-B1	
	Hoff 3	14204-B1	
	Pa	12055-B1	
	Pb	12055-B2	
	PB-1	15692-B3	
	Pc	12055-B3	
	Pf	25102-B1	
	PP7**	15692-B2	
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	Pf3**		J.Virol.1983.47:221-223
	ф-МС		Can.J.Microbiol.1969.15:1179-1186
	Pf1**		J.Mol.Biol.1991.218:349-364
	PR4**		J.Gen.Virol.1979.43:583-592
	A7		J.Bacteriol.1992.174:2407-2411
	KF1		J.Biochem. 1983.93:61-71
	¢CTX**		Mol.Microbiol.1993.4:1703-1709
	f2**		J.Virol.1977.24:135-141
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	φKZ, 21, φNZ,		ddd
	PMN17, PTB80,		
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	PS17**, φ1, 73,		
	M6, Li-2, 7,		
	φmnF82,		
	PTB2, PTB20,		
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	31, PTB21,		
	119x,		
	φPLS27, B3,		
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	Hw12, PM57,	٠.	
	PM62, PM105,		
	148, PM681,		
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	PC131, φC11, SL5,		
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	PM69, PM13,		
	PM61, PM113,		
	φ240, 249 & 269		
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Pseudomonas aeruginosa	297, 309, 318, 11,	Arch.Virol.1993.131:141-151
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	21781-B1	
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PPs-G3	49780-B1	The American Type Culture Collection
Sab 2		Felix d'Herelle Reference
		Centre, Quebec, Quebec
1, 2,3 & 6		Epidemiol.Infect.1995.114:227-236
2a, 3a, 4a, 5a, 6a,		Vet.Med.Nauki.1975.12:55-60
7a, 8a, 9a, 15,		
Epsilon 34		J.Struct.Biol. 1995.115:283-289
	27869-B1	The American Type Culture Collection
	27869-B2	
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		Centre, Quebec, Quebec
	10040_R1	The American Type Culture Collection
	19940-B1	The American Type Culture Collection
Paratumhoid A	19940-B1 12176-B1	The American Type Culture Collection
Paratyphoid A		
Paratyphoid A  Jersey		Felix d'Herelle Reference
Jersey		Felix d'Herelle Reference Centre,Quebec,Quebec
Jersey SasL1, SaL2, Sal		Felix d'Herelle Reference
Jersey SasL1, SaL2, Sal 3,		Felix d'Herelle Reference Centre,Quebec,Quebec
Jersey SasL1, SaL2, Sal 3, SaL4, SaL5 &		Felix d'Herelle Reference Centre,Quebec,Quebec
Jersey SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6	12176-B1	Felix d'Herelle Reference Centre,Quebec,Quebec Indian J.Med.Res. 1997.105:47-52
Jersey SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6 P22**	12176-B1 19585-B1	Felix d'Herelle Reference Centre,Quebec,Quebec
Jersey  SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6  P22** SL-1	12176-B1	Felix d'Herelle Reference Centre,Quebec,Quebec Indian J.Med.Res. 1997.105:47-52  The American Type Culture Collection
Jersey  SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6 P22** SL-1 MB78**	12176-B1 19585-B1	Felix d'Herelle Reference Centre, Quebec, Quebec Indian J.Med.Res. 1997.105:47-52  The American Type Culture Collection  J.Virol. 1982.41: 1038-1043
Jersey SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6 P22** SL-1 MB78** SE1	12176-B1 19585-B1	Felix d'Herelle Reference Centre,Quebec,Quebec Indian J.Med.Res. 1997.105:47-52  The American Type Culture Collection  J.Virol. 1982.41: 1038-1043  J.Gen.Microbiol.1986.132:1035-1041
Jersey  SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6 P22** SL-1 MB78** SE1 LT2	12176-B1 19585-B1	Felix d'Herelle Reference Centre,Quebec,Quebec Indian J.Med.Res. 1997.105:47-52  The American Type Culture Collection  J.Virol. 1982.41: 1038-1043
Jersey SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6 P22** SL-1 MB78** SE1	12176-B1 19585-B1	Felix d'Herelle Reference Centre,Quebec,Quebec Indian J.Med.Res. 1997.105:47-52  The American Type Culture Collection  J.Virol. 1982.41: 1038-1043  J.Gen.Microbiol.1986.132:1035-1041
	1, 2,3 & 6 2a, 3a, 4a, 5a, 6a,	φ6  gh-1  12633-B1  40492-B1  21781-B1

1	Digital Control	T	W-1 C C 1075 100 110 105
	P1CM clr-100	<del> </del>	Mol.Gen.Genet.1975.138:113-126
	F22		Genet.Res.1986.48:139-143
	Fels 1	ļ	J.Gen.Virol.1978.38:263-272
	Fels 2		Genet.Res.1986.48:139-143
	Px		Mol.Gen.Genet.1970.108:184-202
	Plkc		Virology 1974.60:503-514
	A3 & A4		J.Bacteriol. 1987.169:1003-1009
	HT		Genet.Res.1976.27:315-322
Salmonella	IRA		J.Basic Microbiol. 1990.30:707-716
typhimurium	Mudl		Mol.Gen.Genet. 1986.202:327-330
(Cont'd)	P22 (cir4-1, cir5- 1 & cir6-1)		Mol.Gen.Genet.1984.198:105-109
	BF23°°		Mol.Gen.Genet.1976.147:195-202
	Kb1		J.Bacteriol.1974.117:907-908
	P221dis		J.Gen.Virol.1978.41:367-376
	PRD1**		Virology 1990.177:445-451
	I <sub>2</sub> -2**		J.Gen.Microbiol.1982.128:2797-2804
	tf-1		J.Gen.Microbiol.1987.133:953-960
	X**	<u> </u>	J.Gen.Microbiol.1981.126:389-396
Salmonella	8	19937-B1	The American Type Culture Collection
typhosa/typhi	23	19938-B1	3,7
	25	19939-B1	
	46	19942-B1	
	53	19943-B1	
	163	19946-B1	
	175	19947-B1	
	ViI	27870-B1	
	ViVI	27870-B2	
	O1		Felix d'Herelle Refrence Centre, Quebec, Quebec
	ViII		Chung Hua Liu Hsing Ping
	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		H.T.C.1992.13:288
	j2		J.Gen.Microbiol.1983.129:3395-33400
Salmonella sp.	P3	25957-B1	The American Type Culture Collection
	P4**	25957-B2	
	P9a	25957-B3	
	P9c	25957-B4	
	P10	25957-B5	
	102	19945-B1	
	Chi (χ)	9842-B1	
	R34	97541	
	MG40		Virology 1968.34:521-530
	P14		Microb.Pathog.1990.8:393-402
	PSP3		Virology 1992.188:414
	Ike**		Zentralbl.Bakteriol.1976.234:294-304
	P27 & 9NA		J.Virol.1986.12:921-931
Sphaerotilus natans	SNI		Appl.Environ.Microbiol.1979.37:1025-1030

Shigella dysenteriae		23351-B1	The American Type Culture Collection
	P2	11456b	
	<i>∲</i> -80	11456a-B1	
Shigella flexeneri	D20	12661-B1	The American Type Culture Collection
-	SfII**		Mol.Microbiol.1997.26:939-950
	SfV**		Gene 1997.22:217-227
	Sf6**		Mol.Microbiol.1995.18:201-208
	SfX		Gene 1993.129:99-101
Shigella sonnei	C16**		
5g	Ufa		MolBiol (Mosk) 1977.11:323-331
Shigella sp	37	23354-B1	The American Type Culture Collection
Spiroplasma citri	SpV1		Plasmid 1993.29:193-205
Spiroplasma sp.	SpV1-R8A2B		Nucleic Acids Res. 1990.18:1293
<i>Ֆրո օրւաչուա Ֆր.</i>	SpV3		Isr.J.Med.Sci.1987.23:429-433
	Sp V4		J.Bacteriol.1987.169:4950-4961
Staphylococcus albus	Sp +4		Staphylococci & Staphylococcal Infections.1997.
			Vol1:503-508 (Karger,Basel)
			V011.303-308 (184861,24301)
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Staphylococcus aureus		27702-B1	The American Type Culture Collection
		27703-B1	
		27704-B1	
		23360-B1	
		23361-B1	
	15	27705-B1	
	17	27712-B1	
	29	27690-B1	
	42D**	27691-B1	
	42E	27692-B1	
	47	27693-B1	
	52	27694-B1	
	52A	27695-B1	
	53	27696-B1	
	54	27697-B1	
	55	27698-B1	
	71	27699-B1	
	75	27693-B2	
	77	27700-B1	
	79	27701-B1	
	80	27706-B1	
	81	27707-B1	
	83A	27708-B1	
	84	33742	
	85**	33741-B1	
	88	15565	
	92	19685-B1	
	5504'	11987-B1	
	K	11988-B1	
	P1	15752-B1	
İ	P14		
	UC18		

		HER 101 HER 239 HER 283 HER 49	Felix d'Herelle Reference Centre, Quebec, Quebec
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	Twort⁵≎		
	φ11**		J.Bacteriol.1988.170:2409-2411
	φ13°° & φ42°°		J.GenMicrobiol.1989.135:1679-1697
	L54a**		J.Bcteriol.1986.166:385-391
	80α <sup>‡‡</sup>		Can.J.Microbiol.1996.43:612-616
	94,95 & 96		J.Clin.Microbiol.1988.26:2395-2401
	φ131,A <sub>3</sub> & A <sub>5</sub>		Staphylococci & Staphylococcal Infections.1997.
			Vol1:503-508 (Karger,Basel)
	Phi PVL**		Gene 1998.215:57-67
Staphylococcus carnosus	BaSTC2		Felix d'Herelle Reference Centre, Quebec, Quebec
Staphylococcus epidermidis	1a, 2b, 3a, 4b, 5a, 6b, 7b, 8c, 9a, 10a, 11b,12a & 13b		Can.J.Microbiol.1988.34:1358-1361
	41, 63, 118II, 138, 245, 336, 392 & 550		Res.Virol.1994.145:111-121
Staphylococcus	1154A, 1405,		Res. Virol. 1990. 141: 625-635 &
saprophyticus	1314, 1139 & 1259		Res. Virol. 1994. 145:111-121
Staphylococcus sp.	Phi 812, Phi 131, SK311 & U16		Virology 1998.246:241-252
Streptococcus faecalis	VD13	HER44	Felix d'Herelle Reference Centre, Quebec, Quebec
Streptococcus faecium	PE1		Zentralbl.Bakteriol.1975.231:421-425
Streptococcus oralis	Cp-1** & Cp- 7**		FEMS Microbiol.Lett.1989.65:187-192

Streptococcus pneumoniae	Cp-1**	HER223	Felix d'Herelle Reference Centre, Quebec, Quebec
	Cp-1**, Cp-5**,		J.Virol.1981.40:551-559 &
	Cp-7**, Cp-9**,		Eur.J.Biochem.1979.101:59-64 &
	ω-1 & ω-2		Microbial Drug Resistance 1997.3:165-176
	HB-623 & HB-		J.Virol.1990.64:5149-5155
	EJ -1**		J.Bacteriol.1992.174:5516-5525
	Dp-2 & Dp-4		J.Virol.1978.26:221-225
	Dp-1		Virology 1975.63:577-582
	ω-3 & ω-8		J.Virol.1976.19:659-667
	304		J.Bacteriol.1980.141:1298-1304
	HB-1,HB-2, HB-3°°,		J.Bacteriol.1979.138:618-624
	HB-4, HB-5 &		
Street and a series	HB-6	<del> </del>	Mol Missobiology 1007#22-710 720
Streptococcus pyogenes		12202 P1	Mol. Microbiology. 1997#23:719-728
pyogenes	A-1 A-6	12202-B1 12203-B1	The American Type Culture Collection
	A-0 A-25	12203-B1	
	Kjem	14918	
Streptococcus	1	HER 339	Felix d'Herelle Refrence
sp./Enterococcus	182	HER 80	Centre, Quebec, Quebec
7	VD1884	HER 323	
	1A	12169-B1	The American Type Culture Collection
	1B	12170-B1	The financial Type Culture Contonion
	NN	21597-B1	
	42	19948-B1	
	118	19951-B2	
	120	19952-B1	
Veillonella rodentium	N2		Antonie Van Leeuwenhoek 1989.56:263-27
Vibrio cholerae	Psi 92		Intervirology 1993.36:237-244
	VCB-1,2,3 & 4		J.Infetion 1998.36:131
	CP-T1**		J.Virol.1984.51:163-169
	VSK		FEMS Microbiol.Lett.1996.145:17-22
	Phi138		J.Virol.1986.57:960-967
	Phi149		J.Virol.1985.140:217-223
	Fs-2**		Microbiology 1998.144:1901-1906

e4		Felix d'Herelle Reference
		Centre,Quebec,Quebec
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	14100 B1	The American Type Culture Collection
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V	51352-B10	
UTAK		Felix d'Herelle Reference
		Centre, Quebec, Quebec
e <sub>4</sub>		J.Gen.Virol.1987.68:1411-1416
nt1,nt6		Felix d'Herelle Reference
		Centre, Quebec, Quebec
i		Felix d'Herelle Reference Centre, Quebec, Quebec
i .		Centre, Quebec, Quebec
VP1		
ф60		
φHAWI-5		
φPEL8C-1		
α3a		Felix d'Herelle Reference Centre, Quebec, Quebec
NN	11985-B1	The American Type Culture Collection
ł .	51582-B1	
		J.Virol.1987.61:3999-4006
N2		Antonie V.Leeuwenhoek.1989.56:263-271
	e5 X29 β κ 13 14 16 24 32 57 138 145 149 163 N-4 S-5 S-20 M-4 D-10 I II III IV V UTAK e4 nt1,nt6  KVP40** VF33 VP1 φ60 φHAWI-5 φPEL8C-1 α3a NN ph1 Phi149	e5 X29 β κ 13 14 16 24 32 57 138 14100-B1 145 149 14100-B2 149 14100-B30 163 14100-B4 N-4 51352-B1 S-5 51352-B2 S-20 51352-B3 M-4 D-10 51352-B5 I 51352-B6 II 51352-B6 II 51352-B7 III 51352-B8 IV 51352-B9 V 51352-B10 UTAK  e4 nt1,nt6  KVP40** VF33 VP1 φ60 φHAWI-5 φPEL8C-1 α3a  NN 11985-B1 ph1 51582-B1

Yersinia enterocolitica	1		Felix d'Herelle Reference
	2		Centre, Quebec, Quebec
	3		
	4		
	5		
	6		
	7		
	8		
	9		
	φYeO3-12		
	I, IV & VIII		Zentralbl.Bakteriol.Mikrobiol.Hyg.1982.253:1
Yersinia pestis	R	23208-B1	The American Type Culture Collection
•	S	11593-B1	
	Y	23053-B1	
	п		Zh.Mikrobiol.Epidemiol.Immunobiol.1990.11:9
Yersinia pseudotuberculosis	PST**	23207-B1	The American Type Culture Collection
Yersinia sp.	RD2		Mol.Gen.Mikrobiol.Virusol.1990.8:18-21

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Table 2

>Bacteriophage 77, complete genome sequence, 41708 nucleotides

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gatcaaaata cttggggaac ggttagggag taaacttcgc gataatttta aaaattcatg
61
       tataaccccc ctcttataac cattttaagg caggtgatga aatggagatt atagtcgatg
121
       aaaatttagt gottaaagaa aaagaaaggo tacaagtatt atataaagac atacctagca
181
       ataaattaaa agtagttgat ggtttaatta ttcaagcagc aaggctacgt gtaatgcttg
       attacatgtg ggaagacata aaagaaaaag gtgattatga tttatttact caatctgaaa
241
301
       aggegecace atatgaaagg gaaagaccag tagccaaact atttaatgct agagatgctg
       catatcaaaa aataatcaaa caattatcgg atttattgcc cgaagagaaa gaagacacag
361
       aaacgccatc tgatgattac ctatgattag taataaatac gttgatgaat atataaattt
421
481
      gtggaaacaa ggaaagataa ttttaaataa agaaagaatt gatctcttta attatctaca
       aaaacatata tattcacgag atgatgtata ttttgatgaa cagaaaatcg aggattgtat
541
       caaatttatt gaaaaatggt attttccaac attaccattt caaaggttta tcatagctaa
601
       tatatttctt atagataaaa atacagatga agctttcttt acagaatttg ctattttcat
661
721
      gggacgtgga ggcgggaaaa acggtctaat aagtgctatt agtgattttc tttctacgcc
781
       cttacacgga gttaaagaat atcacatctc cattgttgct aatagtgaag atcaagcaaa
      aacatcgttt gatgaaatca gaaccgtttt aatggataac aaacgaaata agacgggtaa
841
901
      aacgccaaaa gctccttatg aagttagtaa agcaaaaata ataaaccgtg caactaaatc
      ggttattcga tataacacat caaacacaaa aaccaaagac ggtggacgtg aggggtgtgt
961
1021
      tatttttgat gaaattcatt atttctttgg tcctgaaatg gtaaacgtca aacgtggtgg
1081
      attaggtaaa aagaaaaata gaagaacgtt ttatataagt actgatggtt ttgttagaga
1141
      gggttatatc gatgcaatga agcacaaaat tgcaagtgta ttaagtggca aggttaaaaa
1201
      tagtagattg tttgcttttt attgtaagtt agacgatcca aaagaagttg atgacagaca
      gacgtgggaa aaggcgaacc caatgttaca taaaccgtta tcagaatacg ctaaaaacact
1261
1321
      gctaagcacg attgaagaag aatataacga tttaccattc aaccgttcaa ataagcccga
1381
      attcatgact aagcgaatga atttgcctga agttgacctt gaaaaagtaa tagcaccatg
      gaaagaaata ctagcgacta atagagagat accaaattta gataatcaaa tgtgtattgg
1441
1501
      tggtttagac tttgcaaaca ttcgagattt tgcaagtgta gggctattat tccgaaaaaa
1561
      cgatgattac atttggttag gacattcgtt tgtaagacaa gggtttttgg atgatgtcaa
1621
      attagaacct cctattaaag aatgggaaaa aatgggatta ttgaccattg tcgatgatga
1681
      tgtcattgaa attgaatata tagttgattg gtttttaaag gctagagaaa aatatgggct
1741
      tgaaaaagtc atagctgata attatagaac tgatattgta agacgtgcgt ttgaggatgc
      tggcataaaa cttgaagtac ttagaaatcc aaaagcaata catggattac ttgcaccacg
1801
      tategataca atgtttgcga aacataacgt aatatatgga gacaatcctt tgatgcgttg
1861
1921
      gtttactaat aatgttgctg taaaaatcaa gccggatgga aataaagagt atatcaaaaa
1981
      agatgaagtc agacgtaaaa cggatggatt catggctttt gttcacgcat tatatagagc
2041
      agacgatata gtagacaaag acatgtctaa agcgcttgat gcattaatga gtatagattt
      ctaatagagg aggtgagaca tgagtattct agaaaagata tttaaaacta ggaaagatat
2101
2161
      aacatatatg cttgatttag atatgataga agatctatca caacaagcgt atgtgaaacg
      tttagcgatt gatagttgta ttgaatttgt tgcgcgagct gtcgctcaaa gtcattttaa
2221
2281
      agtattggaa ggtaatagaa ttcaaaagaa tgatgtttac tacaagttaa atataaaacc
      aaatactgac ttatcaagcg atagtttttg gcaacaagtt atatataaac taatttatga
2341
2401
      taacgaggtt ttaatcgtag taagtgacag caaagaatta cttatcgcag atagctttta
2461
      cagagaagag tacgctttgt atgatgatat attcaaagat gtaacggtta aagattatac
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2521
2581
      gacacacttt gtagaaagtc tattcgaaga ttacgggaaa atattcggaa gaatgatagg
2641
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      cgaaaagaat atagaaaaat tacaagcgtt cacaaataaa ttattcaata cttttaataa
2701
2761
      aaatcaacta gcaatcgcgc ctttgataga aggttttgat tatgaggaat tatctaatgg
      tggtaagaat agtaacatgc ctttttctga attgagtgag ctaatgagag atgcaataaa
2821
2881
      aaatgttgcg ttgatgattg gtatacctcc aggtttgatt tacggagaaa cagctgattt
      ggaaaaaaac acgcttgtat ttgagaagtt ctgtttaaca cctttattaa aaaagattca
2941
3001
      gaacgaatta aacgcgaaac tcataacaca aagcatgtat ttgaaagata caagaataga
3061
      aattgtcggt gtgaataaaa aagacccact tcaatatgct gaagcaattg acaaacttgt
3121
      aagttctggt tcatttacaa ggaatgaggt gcggattatg ttaggtgaag aaccatcaga
      caatcctgaa ttagacgaat acctgattac taaaaactac gaaaaagcta acagtggtga
3181
      aaatgatgaa aaagaaaaag atgaaaacac tttgaaaggt ggtgatgaag atgaaagcgg-
3241
3301
      agattaaagg cgtcatcgtt tccaacgaag ataaatgggt ttacgaaatg cttggtatgg
      attogacttg toctaaagat gttttaacac aactagaatt tagtgatgaa gatgttgata
3361
      ttataattaa ctcaaatggt ggtaacctag tagctggtag tgaaatatat acacatttaa
3421
      gageteataa aggeaaagtg aatgttegta teacageaat ageageaagt geggeatege
3481
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36241 ctaactttat tttaaaaggg cggaaacaat gaaaatcaaa attgaaaaag aaatgaattt 36301 acctgaactt atccaatggg cttgggataa ccccaagtta tcaggtaata aaagattcta 36361 ttcaaatgat gttgagcgca actgttttgt gacttttcat gttgatagca tcttatgtaa 36421 tgtgactgga tatgtatcaa ttaacgataa atttactgtt caagaggaga tataacaatg 36481 aaaatcaaag ttaaaaaaga aatgagatta gatgaattaa ttaaatgggc gcgagaaaat 36541 ccggatctat cacaaggaaa aatattttt tcaacaggat ttagtgatgg attcgttcgt 36601 tttcatccaa atacaaataa gtgttcgacg tcaagtttta ttccaattga tatccccttc 36661 atagttgata ttgaaaaaga agtaacggaa gagactaagg ttgataggtt gattgaatta 36721 ttcgagattc aagaaggaga ctataactct acactatatg agaacactag tataaaagaa 36781 tgtttatatg gcagatgtgt gcctaccaaa gcattctaca tcttaaacga tgacctaact 36841 atgacgttaa tetggaaaga tggggagttg etagtatgat gttgaaattt aaagettggg 36901 ataaagataa aaaagttatg agtattattg acgaaatcga ttttaatagt gggtacattt 36961 tgatttcaac aggttataaa agtttcaatg aagtaaaact attacaatac acaggattta 37021 aagatgtgca cggtgtggag atttatgaag gggatattgt tcaagattgt tattcgagag 37081 aagtaagttt tatcgagttt aaagaaggag ccttttatat aacttttagc aatgtaactg 37141 aattactaag tgaaaatgac gatattattg aaattgttgg aaatattttt gaaaatgaga 37201 tgctattgga ggttatgaga tgacgttcac cttatcagat gaacaatata aaaatctttg 37261 tactaactct aacaagttat tagataaact tcacaaagca ttaaaagatc gtgaagagta 37321 caagaagcaa cgagatgagc ttattgggga tatagcgaag ttacgagatt gtaacaaaga 37381 totagagaag aaagcaagog catgggatag gtattgcaag agogttgaaa aagatttaat 37441 aaacgaattc ggtaacgatg atgaaagagt taaattcgga atggaattaa acaataaaat 37501 ttttatggag gatgacacaa atgaataatc gcgaaaaaat cgaacagtcc gttattagtg 37561 ctagtgcgta taacggtaat gacacagagg ggttgctaaa agagattgag gacgtgtata 37621 agaaagcgca agcgtttgat gaaatacttg agggaatgac aaatgctatt caacattcag 37681 ttaaagaagg tattgaactt gatgaagcag tagggattat ggcaggtcaa gttgtctata 37741 aatatgagga ggaataggaa aatgactaac acattacaag taaaactatt atcaaaaaaat 37801 gctagaatgc ccgaacgaaa tcataagacg gatgcaggtt atgacatatt ctcagctgaa 37861 actgtcgtac tcgaaccaca agaaaaagca gtgatcaaaa cagatgtagc tgtgagtata 37921 ccagagggct atgtcggact attaactagt cgtagtggtg taagtagtaa aacgtattta 37981 gtgattgaaa caggcaagat agacgcggga tatcatggca atttagggat taatatcaag 38041 aatgatgaag aacgtgatgg aataccettt ttatatgatg atatagacge tgaattagaa 38101 gatggattaa taagcattit agatataaaa ggtaactatg tacaagatgg aagaggcata 38161 agaagagttt accaaatcaa caaaggcgat aaactagctc aattggttat cgtgcctata 38221 tggacaccgg aactaaagca agtggaggaa ttcgaaagtg tttcagaacg tggagcaaaa 38281 ggcttcggaa gtagcggagt gtaaagacat cttagatcga gttaaggagg ttttgggggaa 38341 gtgacgcaat acttagtcac aacattcaaa gattcaacag gacgaccaca tgaacatatt 38401 actgtggcta gagataatca gacgtttaca gttattgagg cagagagtaa agaagaagcg 38461 aaagagaagt acgaggcaca agttaaaaga gatgcagtta ttaaagtggg tcagttgtat 38521 gaaaatataa gggagtgtgg gaaatgacgg atgttaaaat taaaactatt tcaggtggag 38581 tttattttgt aaaaacagct gaaccttttg aaaaatatgt tgaaagaatg acgagtttta 38641 atggttatat ttacgcaagt actataatca agaaaccaac gtatattaaa acagatacga 38701 ttgaatcaat cacacttatt gaggagcatg ggaaatgaat cagctgagaa ttttattaca 38761 tgacggtagt agtttgatat tacatgaaga tgaattattt aacgaaatag tatttgtttt 38821 ggacaatttt agaaatgatg atgactattt aacgatagaa aaagattatg gcagagaact 38881 tgtattgaac aaaggttata tagttgggat caatgttgag gaggcagatg atgattaaca 38941 tacctaaaat gaaattcccg aaaaagtaca ctgaaataat caaaaaatat aaaaataaag 39001 cacctgaaga aaaggctaag attgaagatg attttattaa agaaattaaa gataaagaca 39061 gtgaatttta cagtcctacg atggctaata tgaatgaata tgaattaagg gctatgttaa 39121 gaatgatgcc tagtttaatt gatactggag atgacaatga tgattaaaaa acttaaaaat 39181 atggatgggt tegacatett tattgttgga atactgtcat tattcggtat attcgcattg 39241 ctacttgtta tcacattgcc tatctataca gtggctagtt accaacacaa agaattacat 39301 caaggaacta ttacagataa atataacaag agacaagata aagaagacaa gttctatatt 39361 gtattagaca acaaacaagt cattgaaaat tccgacttat tattcaaaaa gaaatttgat 39421 agcgcagata tacaagctag gttaaaagta ggcgataagg tagaagttaa aacaatcggt 39481 tatagaatac actttttaaa tttatatccg gtcttatacg aagtaaagaa ggtagataaa 39541 caatgattaa acaaatacta agactattat tottactago aatgtatgag ttaggtaagt 39601 atgtaactga gcaagtgtat attatgatga cggctaatga tgatgtagag gcgccgagtg 39661 attacgtett tegageggag gtgagtgaat aatgagaata tttatttatg atttgategt 39721 tttgctgttt gctttcttaa tatccatata tattattgat gatggagtga taataaatgc 39781 attaggaatt tttggtatgt ataaaattat agatteettt teagaaaata ttataaagag 39841 gtagataaaa atgaacgagc aaataatagg aagcatatat actttagcag gaggtgttgt 39901 gctttattca gttaaagaga tttttaggta ttttacagat tctaacttac aacgtaaaaa 39961 aatcaattta gaacaaatat atccgatata tttagattgt tttaaaaaagg ctaaaaaagat 40021 gattggagct tatattattc caacaqaaca gcatgaattt ttagattttt ttgatattga 40081 agtotttaat aatttagata agcaaagtaa aaaagogtat gaaaatgtta ttggatttag 40141 acaaatgatt aatttatcaa atagagttaa ggcaatggaa gattttaaga tgagtttcaa 40201 caatgaattt agtacaaatc agattttttt taatccttct tttgttatgg aaacaattgc 40261 tattataaat gaatatcaaa aagatatatc ttatttaaaa aatataatta ataaaatgaa

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40321	tgaaaataga	gcttataatc	atattgatag	ttttatcact	tcagagtacc	gacgaaaaat
40381			ttgataaatt			
40441	aaacagaact	tcgataaaag	aaagaattat	tattaattta	aacaagagga	gatttaaatg
40501	atgtggatta	ctatgactat	tgtatttgct	atattgctat	tagtttgtat	cagtattaat
40561	agtgatcgtg	caagagagat	acaagcactt	agatatatga	atgattatct	acttgatgaa
40621	gtagttaaaa	ctaaagggta	caacgggtta	gaagaataca	ggattgaatt	gaagcgaatg
40681	aataacgata	ttaaaaagta	atttatatta	tcggaggtat	tgcattgaat	gataaagatt
40741	gagaaacacg	atatcaaaaa	gcttgaagaa	tacattcagc	acatcgataa	ctatcgaaga
40801	gagttgaaga	tgcgagaata	tgaattactt	gaaagtcatg	aaccagataa	tgcgggagct
40861	ggcaaaagta	atttgccggg	taacccgatt	gaacgatgtg	caataaagaa	gtttagtgat
40921	aacaggtaca	atacattaag	aaatatagtt	aacggtgtag	atagattgat	aggtgaaagt
40981			attaaggttt			
41041	gaatgggaag	atatagcaca	ttactttggt	acaagtaaga	caagtatatt	acgtagaagg
41101	aatgcactga	tcgataagtt	agcaaagtat	attggttatg	tgtagcggac	ttttacccta
41161	tgtaagtccg	cattaaaaca	gtttattatg	ttagtatcag	attaatattt	aaagttatta
41221	aatgctaata	cgacgcatga	acaagaggcg	catcactatg	tgatgtgtct	ttttatttat
41281	gaggtatgaa	catgttcaaa	ctaattgtaa	atacattact	acacatcaag	tatagatgag
41341	tcttgatact	acttaagtta	tataaggtga	aacattatga	tgactaaaga	cgaacgtata
41401	cgattctata	agtctaaaga	atggcaaata	acaagaaaaa	gagtgctaga	aagagataat
41461	tatgaatgtc	aacaatgtaa	gagagacggc	aagttaacga	catatgacaa	aagcaagcgt
41521	aagtcgttgg	atgtagatca	tatattatcg	ctagaacatc	atccggagtt	tgctcatgac
41581	ttaaacaatt	tagaaacact	gtgtattaaa	tgtcacaaca	aaaaagaaaa	gagatttata
41641	aaaaaagaaa	ataaatggaa	agacgaaaaa	tggtaaatac	ccccgggtca	aaaaaatcaa
41701	aagcgatc					

'

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Table 3

	Name	Position		Name	Position	
1	77ORF005	1957221026	48	77ORF052	17622013	
2	77ORF006	39765196	49	77ORF053	3752137757	
3	77ORF007	2187123076	50	77ORF054	2281823060	
4	77ORF008	21203307	51	77ORF055	1754617788	
5	77ORF009	3194632803	52	77ORF058	1889219122	
6	77ORF010	2609226889	53	77ORF059	3456434785	
7	77ORF011	2444125208	54	77ORF064	2957429795	
8	77ORF012	2978830576	55	77ORF065	2852828746	
9	77ORF013	3362034399	56	77ORF066	2749427703	
10	77ORF014	2776028512	57	77ORF069	3834138547	
11	77ORF015	32914028	58	77ORF070	3626936475	
12	77ORF016	3286733610	59	77ORF071	4049840701	
13	77ORF017	2326923982	60	77ORF072	3873538938	
14	77ORF018	3116931840	61	77ORF073	3094531148	
15	77ORF019	3985140501	62	77ORF074	3854438738	
16	77ORF020	69267570	63	77ORF075	1367313870	
17	77ORF021	3776238304	64	77ORF077	2535725605	
18	77ORF022	3060531156	65	77ORF079	2908929280	
19	77ORF023	2690327346	66	77ORF080	3520435389	
20	77ORF024	1070011140	67	77ORF085	2406024242	
21	77ORF025	970710147	68	77ORF092	3970639876	
22	77ORF026	4072941145	69	77ORF094	3222632393	
23	77ORF027	65186925	70	77ORF096	1360613773	
24	77ORF028	3479535199	71	77ORF098	70927256	
25	77ORF029	61176521	72	77ORF102	2905129212	
26	77ORF030	3647836879	73	77ORF104	3439334551	
27	77ORF031	3915139546	74	77ORF109	1828218434	
28	77ORF032	3389234266	75	77ORF112	3954339692	
29	77ORF033	57586120	76	77ORF117	2736127501	
30	77ORF034	78868236	77	77ORF118	3839038530	
31	77ORF035	1925819560	78	77ORF120	3605936199	
32	77ORF036	3687637223	79	77ORF124	3369933833	
33	77ORF037	102446	80	77ORF128	1422114355	
34	77ORF038	3490835219	81	77ORF130	1567515806	
35	77ORF039	3722037528	82	77ORF133	84148542	
36	77ORF040	4137741676	83	77ORF140	1311313235	
37	77ORF041	3545435753	84	77ORF147	70297148	
38	77ORF042	54905774	85	77ORF149	3066830787	
39	77ORF043	2930429564	86	77ORF151	3183731953	
40	77ORF044	1848118768	87	77ORF155	3027830391	
41	77ORF045	52165500	88	77ORF157	40444157	
42	77ORF046	2566325935	89	77ORF167	2069220799	
43	77ORF047	1115911425	90	77ORF175	3571735821	
44	77ORF048	2877629039	91	77ORF176	68366940	
45	77ORF049	3601336255	92	77ORF178	3539035491	
46	77ORF050	3575336007	93	770RF179	83188419	<b>-</b>
47	77ORF051	3893139167	94	77ORF182	2926829564	

## Table 4

## 770RF017 sequence

23982		ato	gacg	rcat	aat	ata	gaa	aaa	.cgc	att	aat	aaa	tta	aaaa	cttct
1 M	T	H	N	I	E	K	R	I	N	K	L	K	T	S	
23937		gga	aat	сса	aaa	ttt	aaa	aag	tta	gat	tca	gat	att	cact	attta
16 G	N	P	K	F	K	K	L	D	S	D	I	H	Y	L	
23892		cto	aag	aga	ttt	gaa	ggt	gaa	aaa	aac	cat	aaa	ggt	tttt	atcca
31 L	K		F										Y		
23847		aag	gttt	aaa	caa	gga	gaa	ata	gtt	ttt	gta	gat	ttc	ggta	taaac
46 K	F	K	Q	G	E	I	V	F	V	D	F	G	I	N	
23802		gtt	aat	aaa	gaa	ttt	tct	aat	tca	cac	ttt	gca	ata	gtga	ıtgaat
61 V	N		E	F					_	Α		V			
23757		aaa													cctta
76 K	N	D	S	N	T	E	D	I	V	N	V	I	P	L	
23712		tcc	ctct											tttg	atttg
91 S	S	K	E								N				
23667		aaa	itgg	gag	tat	tat	tta	aga	ttg	ttt	tta	aat	tta	atta	gcgcg
106 K	W	E		Y				F			L	I	S	A	
23622		caa	aat	aat	tca	gct	ata	tta	aaa	gaa	gtt	ttc	gat	aaaa	aatac
23622 121 Q	N	N	S	Α	I	L	K	E	V	F	D	K	K	Y	
	N	N	S	Α	I	L	K gaa	E ttc	V atc	F act	D aaa	K gat	K tat	Y	aatac
121 Q		N	S	Α	I	L	K gaa	E ttc	V atc	F act	D	K gat	K tat	Y ttta	
121 Q 23577		N caa N	S aaaa N	A .aac T	I aac E	L aca F	K gaa I	E ttc T	V atc K	F act D	D aaa Y	K gat F	K tat I	Y ttta E	
121 Q 23577 136 Q		N caa N ttt	S aaaa N ata D	A .aac T .tct S	I aac E gat L	L aca F agt E	K gaa I tta I	E ttc T gaa E	V atc K att N	F act D gaa K	D aaa Y aat L	K gat F aaa N	K tat I tta K	Y ttta E aata I	ittgaa laaatt
121 Q 23577 136 Q 23532	K	N caa N ttt	S aaaa N ata D	A .aac T .tct S	I aac E gat L att	L aca F agt E aat	K gaa I tta I aac	E ttc T gaa E ata	V atc K att N gta	F act D gaa K tca	D aaa Y aat L gca	K gat F aaa N att	K tat I tta K gat	Y ttta E aata I aagg	ittgaa
121 Q 23577 136 Q 23532 151 F	K	N caa N ttt S gao	S Naaa N ata D aga	A aac tct S aac	I aac E gat L att	L aca F agt E aat	K gaa I tta I aac V	E ttc T gaa E ata S	V atc K att N gta A	F act D gaa K tca	D aaa Y aat L gca D	K gat F aaa N att K	K tat I tta K gat V	Y ttta E aata I aagg K	attgaa aaatt gtaaaa
121 Q 23577 136 Q 23532 151 F 23487	K	N caa N ttt S gao	S Naaa N ata D aga	A aac T s tct S aac	I aac gat L att N	L aca F agt E aat aat	K gaa I tta I aac V agt	E ttc gaa E ata stac	V atc K att N gta A gct	F act gaa K tca tgc	D aaa Y aat L gca D ata	K gat aaa N att aat	K tat I tta K gat V tct	Y ttta E aata I aagg K	ittgaa laaatt
121 Q 23577 136 Q 23532 151 F 23487 166 D	K I R	N caa N ttt	S Naaa N Cata D Caga I Itta G	A .aac .tct S .aac N	I aac gat L att N ggt	L aca F agt E aat aat	K gaa I tta I aac V agt	E ttc gaa E ata stac	V atc K att N gta A gct	F act gaa K tca tgc N	D aaa Y aat L gca D ata	K gat aaa N att att aat	K tat I tta K gat V tct Q	Y ttta E aata I aagg K ttcc	attgaa aaatt taaaa agccg
121 Q 23577 136 Q 23532 151 F 23487 166 D 23442	K I R	N caa N ttt	S Naaa N Cata D Caga I Itta G	A .aac .tct S .aac N	I aac gat L att N ggt	L aca F agt E aat aat	K gaa I tta I aac V agt A	E ttc gaa E ata stac tac	V atc K att N gta A gct aaa	F act gaa K tca tgc N	D aaa Y aat L gca D ata S tta	K gat aaa N att k aat ccc	K tat I tta K gat V tct Q	Y ttta E aata I aagg K ttcc	attgaa aaatt gtaaaa
121 Q 23577 136 Q 23532 151 F 23487 166 D 23442 181 K	K I R	N caa N ttt S gac N aaa K att K	S aaaa N aata D aaga I atta G agt	A .aac T .s.aac .aac N .aaa N aag	I aac gat att agg ttt I	aca F agt E aat aat cgc	K gaa I tta I aac V agt ata K	E ttc gaa E ata s tac aga V	V atc K att Sta G A G C I aaa L	Fact Daa Ktca tca tgc ytt	D aaa Y aat C C C C C C C C C C C C C C C C C C	K gat aaa N att aat ccc K	K tat I tta K gat tct Caa I	Y ttta E aata I aagg K ttcc P aaaa	attgaa aaatt taaaa agccg
121 Q 23577 136 Q 23532 151 F 23487 166 D 23442 181 K 23397	K I R L	N caa N ttt S gac N aaa K att K	S aaaa N aata D aaga I atta G agt	A .aac T .s.aac .aac N .aaa N aag	I aac gat att agg ttt I	aca F agt E aat aat cgc	K gaa I tta I aac V agt ata K	E ttc gaa E ata s tac aga V	V atc K att Sta G A G C I aaa L	Fact Daa Ktca tca tgc ytt	D aaa Y aat C C C C C C C C C C C C C C C C C C	K gat aaa N att aat ccc K	K tat I tta K gat tct Caa I	Y ttta E aata I aagg K ttcc P aaaa	attgaa aaatt taaaa agccg
121 Q 23577 136 Q 23532 151 F 23487 166 D 23442 181 K 23397 196 I	K I R L	N caa N ttt S gao N aaa K att K aat	S aaaa N ata D aga Itta G agt cca I	A aac tct saac aaa n aag agta D	I aac gat att ata	L aca F t agt agt agt ag I t c R t c R t c R t	K gaa tta a V ta A A tct	E ttc gaa stac aga tcg	V atc att gtA gtA gtaa gat gat	F act gaa tca tgc gtt att	D aaa Y aa L a C C C C C C C C C C C C C C C C	K gat aaa N a K t a K t C K t t t I	K tat I tta gat Caa Ctg	Y ttta E aata I aagg K ttcc P aaaa K ataa R	attgaa aaatt gtaaaa agccg attaaa ataga
121 Q 23577 136 Q 23532 151 F 23487 166 D 23442 181 K 23397 196 I 23352	K I R L	N caa N ttt S gao N aaa K att K aat	S aaaa N ata D aga Itta G agt cca I	A aac tct saac aaa n aag agta D	I aac gat att ata	L aca F t agt agt agt ag I t c R t c R t c R t	K gaa tta a V ta A A tct	E ttc gaa stac aga tcg	V atc att gtA gtA gtaa gat gat	F act gak at to get at to get at to get at to get at to get at to constitution of the second	D aaa Y aa L a C C C C C C C C C C C C C C C C	Kat aa N t aa F C K t I a	K tat I tta gat Caa Ctg	Y ttta E aata I aagg K ttcc P aaaa K ataa R	attgaa aaatt taaaa agccg

# Physico-chemical parameters of ORF 770RF017

1	MTHNIEKRIN	KLKTSGNPKF	KKLDSDIHYL	LKRFEGEKNH	KGFYPKFKQG	EIVFVDFGIN
61	VNKEFSNSHF	AIVMNKNDSN	TEDIVNVIPL	SSKENKKYLK	MNFDLKWEYY	LRLFLNLISA
121	QNNSAILKEV	FDKKYQKNNT	EFITKDYFIE	FISDSLEIEN	KLNKIDRNIN	NIVSAIDKVK
181	KLKGNSYACI	NSFQPISKFR	IRKVLPQKIK	NPVIDSSDIM	LLINRINNNI	LQIPDIR

Number of amino acids:	237
Average molecular weight (Daltons):	27887.38
Mean amino acid weight (Daltons):	117.67
Monoisotopic molecular weight (Daltons):	27869.83
Mean amino acid monoisotopic weight (Daltons):	117.59

## Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	5	2.11%	7.58%	Cys	С	1	0.42%	1.66%
Asp	D	14	5.91%	5.28%	Glu	E	13	5.49%	6.37%
Phe	F	16	6.75%	4.09%	Gly	G	6	2.53%	6.84%
His	Н	4	1.69%	2.24%	Ile	I	29	12.24 %	5.81%
Lys	K	33	13.92 %	5.95%	Leu	L	19	8.02%	9.42%
Met	М	4	1.69%	2.37%	Asn	N	30	12.66 %	4.45%
Pro	P	7	2.95%	4.9%	Gln	Q	6	2.53%	3.97%
Arg	R	8	3.38%	5.16%	Ser	S	17	7.17%	7.12%
Thr	T	5	2.11%	5.67%	Val	V	11	4.64%	6.58%
Trp	W	1	0.42%	1.23%	Tyr	Y	8	3.38%	3.18%

Number of acidic (negative) amino acids (ED):	27
	11.39%
Number of basic (positive) amino acids (KR):	41
•	17.30%
Total charge (KRED):	68
•	28.69%
Net charge (KR - ED):	14
	5.91%
Theoritical pI:	10.01
Total linear charge density:	0.30
Average hydrophobicity:	-5.37
Ratio of hydrophilicity to hydrophobicity:	1.41
Percentage of hydrophilic amino acid:	57.81%
Percentage of hydrophobic amino acid:	42.19%
Ratio of %hydrophilic to %hydrophobic:	1.37

155

#### 77ORF019 sequence

atgaacgagcaaataataggaagcatatatactttagcaggaggt 1 MNEQIIGSIYTLAGG 39896 gttgtgctttattcagttaaagagatttttaggtattttacagat 16 V V L Y S V K E I F R Y F T D 39941 tctaacttacaacgtaaaaaaatcaatttagaacaaatatatccq 31 SNLORKKINLEQIYP 39986 atatatttagattgttttaaaaaggctaaaaagatgattggagct 46 IYLDCFKKAKKMIGA tatattattccaacagaacagcatgaatttttagattttttgat 40031 61 Y I I P T E Q H E F L D F F D attgaagtctttaataatttagataagcaaagtaaaaaagcgtat 40076 76 IEVFNNLDKQSKKAY gaaaatgttattggatttagacaaatgattaatttatcaaataga 40121 91 ENVIGFRQMINLSNR gttaaggcaatggaagattttaagatgagtttcaacaatgaattt 40166 106 V K A M E D F K M S F N N E F 40211 agtacaaatcagattttttttaatccttcttttgttatggaaaca 121 S T N Q I F F N P S F V M E T attgctattataaatgaatatcaaaaagatatatcttatttaaaa 40256 136 I A I I N E Y Q K D I S Y L K aatataattaataaaatgaatgaaaatagagcttataatcatatt 40301 151 N I I N K M N E N R A Y N H I gatagttttatcacttcagagtaccgacgaaaaataaacgattat 40346 166 D S F I T S E Y R R K I N D Y aatctttatcttgataaatttgaagaacagtttagtcaaaagttt 40391 181 N L Y L D K F E E Q F S Q K F 40436 aaaataaacagaacttcgataaaagaaagaattattattaattta 196 K I N R T S I K E R I I I N L aacaagaggagatttaaatga 40501 40481 211 N K R R F K \*

# Physico-chemical parameters of ORF 77ORF019

1	MNEQIIGSIY	TLAGGVVLYS	VKEIFRYFTD	SNLQRKKINL	EQIYPIYLDC	FKKAKKMIGA
61	YIIPTEQHEF	LDFFDIEVFN	NLDKQSKKAY	ENVIGFRQMI	NLSNRVKAME	DFKMSFNNEF
121	STNQIFFNPS	<b>FVMETIAIIN</b>	EYQKDISYLK	NIINKMNENR	AYNHIDSFIT	SEYRRKINDY
181	NLYLDKFEEQ	FSOKFKINRT	SIKERIIINL	NKRRFK		

Number of amino acids:	216
Average molecular weight (Daltons):	26026.06
Mean amino acid weight (Daltons):	120.49
Monoisotopic molecular weight (Daltons):	26009.34
Mean amino acid monoisotopic weight (Daltons):	120.41

## Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb %		Average % in Swissprot	
Ala	A	7	3.24%	7.58%	Cys	С	1	0.46%	1.66%	
Asp	D	10	4.63%	5.28%	Glu	E	16	7.41%	6.37%	
Phe	F	19	8.80%	4.09%	Gly	G	5	2.31%	6.84%	
His	Н	2	0.93%	2.24%	Ile	I	28	12.96 %	5.81%	
Lys	K	22	10.19 %	5.95%	Leu	L	12	5.56%	9.42%	
Met	М	7	3.24%	2.37%	Asn	N	23	10.65 %	4.45%	
Pro	P	3	1.39%	4.9%	Gln	Q	10	4.63%	3.97%	
Arg	R	11	5.09%	5.16%	Ser	S	13	6.02%	7.12%	
Thr	Т	7	3.24%	5.67%	Val	V	7	3.24%	6.58%	
Trp	W	0	0.00%	1.23%	Tyr	Y	13	6.02%	3.18%	

26
12.04%
33
15.28%
59
27.31%
7
3.24%
9.52
0.28
-4.84
1.37
54.17%
45.83%
1.18

### 77ORF043 sequence

293	04		atg	tat	tac	gaa	ata	ggc	gaa	atc	ata	.cgc	aaa	aat	attca	atgtt
1	M	Y	Y	E	I	G	E	I	I	R	K	N	I	H	V	
293	49		aac	gga	ttc	gat	ttt	aag	cta	ttc	att	tta	aaa	ggt	catai	gggc
16	N	G	F	D	F	K	L	F	I	L	K	G	H	M	G	
293	94		ata	tca	ata	caa	gtt	aaa	gat	atg	aac	aac	gta	.cca	attaa	aacat
31	I	S	I	Q	V	K	D	М	N	N	V	P	I	K	H	
294	39		gct	tat	gto	gta	gat	gag	aat	gac	tta	gat	atg	gca	tcaga	actta
46	Α	Y	V	. V	D	E	N	D	L	D	M	A	S	D	L	
294	84		ttt	aac	caa	gca	ata	gat	gaa	tgg	att	gaa	gag	aac	acaga	acgaa
61	F	N	Q	Α	I	D	E	W	I	E	Ε	N	Т	D	E	
295	29		cag	gac	aga	cta	att	aac	tta	gtc	atg	aaa	tgg	tag	295	54
76	Q	D	R	L	I	N	L	V	M	K	W	*				

# Physico-chemical parameters of ORF 77ORF043

1 MYYEIGEIIR KNIHVNGFDF KLFILKGHMG ISIQVKDMNN VPIKHAYVVD ENDLDMASDL

61 FNQAIDEWIE ENTDEQDRLI NLVMKW

Number of amino acids:

Average molecular weight (Daltons):

Mean amino acid weight (Daltons):

Monoisotopic molecular weight (Daltons):

Mean amino acid monoisotopic weight (Daltons):

10180.02

Mean amino acid monoisotopic weight (Daltons):

118.37

### Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	3	3.49%	7.58%	Cys	C	0	0.00%	1.66%
Asp	D	9	10.47 %	5.28%	Glu	Е	7	8.14%	6.37%
Phe	F	4	4.65%	4.09%	Gly	G	4	4.65%	6.84%
His	Н	3	3.49%	2.24%	Ile	I	11	12.79 %	5.81%
Lys	K	6	6.98%	5.95%	Leu	L	6	6.98%	9.42%
	M	5	5.81%	2.37%	Asn	N	8	9.30%	4.45%
Pro	P	1	1.16%	4.9%	Gln	Q	3	3.49%	3.97%
Arg	R	2	2.33%	5.16%	Ser	S	2	2.33%	7.12%
Thr	T	1	1.16%	5.67%	Val	V	6	6.98%	6.58%
Trp	W	2	2.33%	1.23%	Tyr	Y	3	3.49%	3.18%

Number of acidic (negative) amino acids (ED):	16 18.60%
Number of basic (positive) amino acids (KR):	8 9.30%
Total charge (KRED):	24 27.91%
Net charge (KR - ED): 9.30%	-8 -
Theoritical pI: Total linear charge density:	4.38 0.30
Average hydrophobicity:	-2.80 1.19
Ratio of hydrophilicity to hydrophobicity: Percentage of hydrophilic amino acid:	48.84%
Percentage of hydrophobic amino acid: Ratio of %hydrophilic to %hydrophobic:	51.16% 0.95

#### 77ORF102 sequence

29051 atgageacatttataaaagetacctagtageagtattatgette

1 M S N I Y K S Y L V A V L C F

29096 acagtettagegattgtacttatgeegtttetatactteactaca

16 T V L A I V L M P F L Y F T T

29141 gcatggteaattgegggattegeaagtategeaacatteatgtac

31 A W S I A G F A S I A T F M Y

29186 tacaaagaatgettttteaaagaataa 29212

46 Y K E C F F K E \*

# Physico-chemical parameters of ORF 77ORF102

1 MSNIYKSYLV AVLCFTVLAI VLMPFLYFTT AWSIAGFASI ATFMYYKECF FKE

Number of amino acids:	53
Average molecular weight (Daltons):	6155.42
Mean amino acid weight (Daltons):	116.14
Monoisotopic molecular weight (Daltons):	6151.07
Mean amino acid monoisotopic weight (Daltons):	116.06

## Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo I	Numb er	%	Average % in Swissprot
Ala	Α	6	11.32 %	7.58%	Cys	С	2	3.77 %	1.66%
Asp	D	0	0.00%	5.28%	Glu	Е	2	3.77 %	6.37%
Phe	F	7	13.21 %	4.09%	Gly	G	1	1.89 %	6.84%
His	Н	0	0.00%	2.24%	Ile	I	4	7.55 %	5.81%
Lys	K	3	5.66%	5.95%	Leu	L	5	9.43 %	9.42%
Met	М	3	5.66%	2.37%	Asn	N	1	1.89 %	4.45%
Pro	Р	1	1.89%	4.9%	Gln	Q	0	0.00 %	3.97%
Arg	R	0	0.00%	5.16%	Ser	s	4	7.55 %	7.12%
Thr	Т	4	7.55%	5.67%	Val	v	4	7.55 %	6.58%
Trp	w	1	1.89%	1.23%	Tyr	Y	5	9.43 %	3.18%

Number of acidic (negative) amino acids (ED):	2
, ,	3.77%
Number of basic (positive) amino acids (KR):	3
<u>-</u>	5.66%
Total charge (KRED):	5
-	9.43%
Net charge (KR - ED):	1
	1.89%
Theoritical pI:	8.18
Total linear charge density:	0.13
Average hydrophobicity:	10.81
Ratio of hydrophilicity to hydrophobicity:	0.40
Percentage of hydrophilic amino acid:	28.30%
Percentage of hydrophobic amino acid:	71.70%

WO 00/32825

161

Ratio of %hydrophilic to %hydrophobic:

0.39

#### 77ORF104 sequence

atggtaaccaaagaatttttaaaaactaaacttgagtgttcagat

M V T K E F L K T K L E C S D

34438 atgtacgctcagaactcatagatgaggcacagggcgatgaaaat

16 M Y A Q K L I D E A Q G D E N

34483 aggttgtacgacctatttatccaaaaacttgcagaacgtcataca

31 R L Y D L F I Q K L A E R H T

34528 cgccccgctatcgtcgaatattaa 34551

46 R P A I V E Y \*

# Physico-chemical parameters of ORF 77ORF104

1 MVTKEFLKTK LECSDMYAQK LIDEAQGDEN RLYDLFIQKL AERHTRPAIV EY

Number of amino acids:	52
Average molecular weight (Daltons):	6193.13
Mean amino acid weight (Daltons):	119.10
Monoisotopic molecular weight (Daltons):	6189.12
Mean amino acid monoisotopic weight (Daltons):	119.02

### Amino acid composition

Aci d	Symbo I	Numb er	%	Average % in Swissprot	Aci d	Symbo 1	Numb er	%	Average % in Swissprot
Ala	A	4	7.69 %	7.58%	Cys	С	1	1.92%	1.66%
Asp	D	4	7.69 %	5.28%	Glu	E	6	11.54 %	6.37%
Phe	F	2	3.85 %	4.09%	Gly	G	1	1.92%	6.84%
His	Н	1	1.92 %	2.24%	Ile	I	3	5.77%	5.81%
Lys	K	5	9.62 %	5.95%	Leu	L	6	11.54 %	9.42%
Met	М	2	3.85 %	2.37%	Asn	N	1	1.92%	4.45%
Pro	P	1	1.92 %	4.9%	Gln	Q	3	5.77%	3.97%
Arg	R	3	5.77 %	5.16%	Ser	S	1	1.92%	7.12%
Thr	Т	3	5.77 %	5.67%	Val	V	2	3.85%	6.58%
Тгр	w	0	0.00 %	1.23%	Tyr	Y	3	5.77%	3.18%

Number of acidic (negative) amino acids (ED):	10
Number of basic (positive) amino acids (KR):	19.23% 8
Transport of basis (Postario)	15.38%
Total charge (KRED):	18 34.62%
Net charge (KR - ED):	-2 -
3.85%	5.02
Theoritical pI:	5.03
Total linear charge density:	0.38
Average hydrophobicity:	-5.81
Ratio of hydrophilicity to hydrophobicity:	1.47
Percentage of hydrophilic amino acid:	53.85%
Percentage of hydrophobic amino acid:	46.15%

164

Ratio of %hydrophilic to %hydrophobic:

1.17

PCT/IB99/02040

## 77ORF182 sequence

2926	8		atg	ttc	aat	ata	aaa	cga	aaa	acg	gag	gaa	gtc	aag	atgi	tattac
1	М	F	N	I	K	R	K	T	E	Ε	V	K	M	Y	Y	
2931	.3		gaa	ata	ggc	gaa	atc	ata	cgc	aaa	aat	att	cat	gtt	aac	ggattc
16	Е	I	Ğ	E	Ī	Ī	R	K	N	I	Н	V	N	G	F	
2935	8		gat	ttt	aag	cta	ttc	att	tta	aaa	ggt	cat	atg	ggc	ata	tcaata
31	D	F	ĸ	L	F	I	L	K	G	H	M	G	I	S	I	
2940	3		caa	gtt	aaa	gat	atg	aac	aac	gta	cca	att	aaa	cat	gcti	tatgtc
46	0	V	K	D	M	N	N	V	P			H	Α		V	
2944	8		gta	gat	gag	aat	gac	tta	gat	atg	gca	tca	gac	tta	ttt	aaccaa
61	V	D	E	N	֓֞֞֞֞	L	D		Α		D	L	F	N	Q	
2949	3		gca	ata	gat	gaa	tgg	att	gaa	gag	aac	aca	gac	gaa	cag	gacaga
76	Α	I	D	E	W	I	E	E	N	$\mathbf{T}$	D	E	Q	D	R	
2953	8		cta	att	aac	tta	gto	atg	aaa	tgg	tag	29	564			
91	L	I	N	L	V	M	K	W	*							

# Physico-chemical parameters of ORF 77ORF182

1 MFNIKRKTEE VKMYYEIGEI IRKNIHVNGF DFKLFILKGH MGISIQVKDM NNVPIKHAYV

61 VDENDLDMAS DLFNQAIDEW IEENTDEQDR LINLVMKW

Number of amino acids:98Average molecular weight (Daltons):11691.50Mean amino acid weight (Daltons):119.30Monoisotopic molecular weight (Daltons):11683.84Mean amino acid monoisotopic weight (Daltons):119.22

#### Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	3	3.06 %	7.58%	Cys	С	0	0.00%	1.66%
Asp	D	9	9.18 %	5.28%	Glu	E	9	9.18%	6.37%
Phe	F	5	5.10 %	4.09%	Gly	G	4	4.08%	6.84%
His	Н	3	3.06 %	2.24%	Ile	I	12	12.24 %	5.81%
Lys	K	9	9.18 %	5.95%	Leu	L	6	6.12%	9.42%
Met	М	6	6.12 %	2.37%	Asn	N	9	9.18%	4.45%
Pro	P	1	1.02 %	4.9%	Gln	Q	3	3.06%	3.97%
Arg	R	3	3.06 %	5.16%	Ser	S	2	2.04%	7.12%
Thr	Т	2	2.04 %	5.67%	Val	V	7	7.14%	6.58%
Ттр	w	2	2.04 %	1.23%	Tyr	Y	3	3.06%	3.18%

Number of acidic (negative) amino acids (ED):	18
	18.37%
Number of basic (positive) amino acids (KR):	12
•	12.24%
Total charge (KRED):	30
• .	30.61%
Net charge (KR - ED):	-6 -
6.12%	en en en en en en en en en en en en en e
Theoritical pI:	4.76
Total linear charge density:	0.33
Average hydrophobicity:	-3.89
Ratio of hydrophilicity to hydrophobicity:	1.28

Percentage of hydrophilic amino acid:	-	51.02%
Percentage of hydrophobic amino acid:		48.98%
Ratio of %hydrophilic to %hydrophobic:		1.04

#### Table 5

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100017|lan|770RF017 Phage 77 ORF |23269-23982|-3 (237 letters)

Database: nr

393,678 sequences; 120,452,765 total letters

	Score	E
Sequences producing significant alignments:	(bits)	Value
gi 4493986 emb CAB39045.1  (AL034559) predicted using hexExon;	. 41	0.010
gi 730607 sp P23250 RPI1 YEAST NEGATIVE RAS PROTEIN REGULATOR P	. 38	0.053
gi 3097044 emb CAA75299  (Y15035) K1R [Cowpox virus]	38	0.090
gi 2146245 pir   S73794 hypothetical protein H91_orf180 - Mycopl	. 38	0.090
gi 83910 pir  S04682 ribosomal protein varl - yeast (Candida gl	. 37	0.15
gi 133135 sp P21358 RMAR_CANGA MITOCHONDRIAL RIBOSOMAL PROTEIN	. 37	0.15
gi 2128843 pir  H64475 hypothetical protein MJ1409 - Methanococ	. 36	0.20
gi 5107017 gb AAD39926.1 AF126285_2 (AF126285) RNA polymerase [	. 36	0.35
qi 2146210 pir  \$73342 hypothetical protein E07_orf166 - Mycopl	. 35	0.60

Database: swissprot

79,449 sequences; 28,874,452 total letters

Seq	uences j	producing si	gnificant alignments:	Score (bits)	E Value
sp	P23250	RPI1 YEAST	NEGATIVE RAS PROTEIN REGULATOR PROTEIN.	38	0.014
sp	P21358	RMAR CANGA	MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	37	0.040
sp	Q21444	LDLC CAEEL	LDLC PROTEIN HOMOLOG.	34	0.35
sp	P27240	RFAY ECOLI	LIPOPOLYSACCHARIDE CORE BIOSYNTHESIS PROT.	. 33	0.46
sp	P53192	YGCO YEAST	HYPOTHETICAL 27.1 KD PROTEIN IN ALK1-CKB1.	. 33	0.60
sp	P32908	SMC1 YEAST	CHROMOSOME SEGREGATION PROTEIN SMC1 (DA-B.	. 33	0.60
ge	P54683	TAGE DICDI	PRESTALK-SPECIFIC PROTEIN TAGB PRECURSOR .		0.78
sp	Q03100	CYAA DICDI	ADENYLATE CYCLASE, AGGREGATION SPECIFIC (.	. 32	0.78

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BLASTP 2.0.8 [Jan-05-1999]
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Query= sid|100019|lan|770RF019 Phage 77 ORF|39851-40501|2 (216 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

	Score	E
Sequences producing significant alignments:	bits)	Value
qi 3341966 dbj BAA31932  (AB009866) orf 59 [bacteriophage phi PVL	] 437	e-122
gil2689911 (AE000792) B. burgdorferi predicted coding region BB	. 38	0.058
gi 1171589 emb CAA64574 (X95275) frameshift (Plasmodium falcip	. 37	0.10
gi 4493986 emb CAB39045.1 (AL034559) predicted using hexExon;	. 36	0.23
gi 141257 sp P18019 YPI9 CLOPE HYPOTHETICAL 14.5 KD PROTEIN (OR	. 36	0.29
gi 133412 sp P27059 RPOB ASTLO DNA-DIRECTED RNA POLYMERASE BETA	. 35	0.51
gi 3122231 sp Q58851 HISX_METJA HISTIDINOL DEHYDROGENASE (HDH)	. 35	0.51
gi 3649757 emb CAB11106.1 (Z98547) predicted using hexExon; MA	. 34	0.66
gi 2688313 (AE001146) sensory transduction histidine kinase, pu	. 34	0.87

Database: swissprot

79,449 sequences; 28,874,452 total letters

Sequences	producing s	significant alignments:	Score (bits)	E Value
sp   P18019	YPI9_CLOPE	HYPOTHETICAL 14.5 KD PROTEIN (ORF9).	36	0.079
	HISX METJA		35	0.14
	RPOB ASTLO		35	0.14
sp   002224	CENE HUMAN	CENTROMERIC PROTEIN E (CENP-E PROTEIN).	34	0.31
	ARP PLAFA	ASPARAGINE-RICH PROTEIN (AG319) (ARP) (FRA.	. 33	0.53
	IPAB SHIFL	62 KD MEMBRANE ANTIGEN.	32	0.69
	VTA2 XENLA	VITELLOGENIN A2 PRECURSOR (VTG A2) [CONTA.	. 32	0.90
	CP3H_CAVPO	CYTOCHROME P450 3A17 (EC 1.14.14.1) (CYPI	. 32	0.90
	RMAR_CANGA	MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	32	0.90
	IPAB SHIDY	62 KD MEMBRANE ANTIGEN.	32	1.2

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100043|lan|770RF043 Phage 77 ORF|29304-29564|3 (86 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

-	core its)	E Value
gi 3341947 dbj BAA31913  (AB009866) orf 39 [bacteriophage phi PVL] gi 744518 prf  2014422A FKBP-rapamycin-associated protein [Homo gi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN gi 1169735 sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTE gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa gi 3875402 emb CAA98122  (Z73906) cDNA EST EMBL:D64544 comes fr gi 1084792 pir  S54091 hypothetical protein YPR070w - yeast (Sa	32 32 32 32 31	0.84 0.84 0.84 0.84 2.5

Database: swissprot

79,449 sequences; 28,874,452 total letters

		Score	E
Sequences	producing significant alignments:	(bits)	Value
sp   P42345	FRAP HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP)	. 32	0.24
- !	FRAP RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R		0.24
sp P34554	YNP1 CAEEL HYPOTHETICAL 42.2 KD PROTEIN T05G5.1 IN C	. 28	3.5
sp Q24118	LIO DROME LINOTTE PROTEIN.	28	3.5
sp P80034	ACH2_BOMMO ANTICHYMOTRYPSIN II (ACHY-II).	28	3.5
sp   P22922	Alat_BOMMO ANTITRYPSIN PRECURSOR (AT).	28	3.5
sp Q44363	TRAA AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	28	3.5
sp P38255	YBUS YEAST HYPOTHETICAL 51.3 KD PROTEIN IN PHO5-VPS1	. 27	6.0
sp   P55822	SH3B_HUMAN SH3BGR PROTEIN (21-GLUTAMIC ACID-RICH PRO	. 27	7.9
sp   Q58482	YA82 METJA HYPOTHETICAL PROTEIN MJ1082.	27	7.9
sp P34252	YKK8 YEAST HYPOTHETICAL 52.3 KD PROTEIN IN HAP4-AAT1	. 27	7.9

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100102|lan|770RF102 Phage 77 ORF|29051-29212|2 (53 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 3341946 dbj BAA31912  (AB009866) orf 38 [bacteriophage phi P gi 4325288 gb AAD17315  (AF123593) voltage-dependent sodium cha gi 2649684 (AE001040) A. fulgidus predicted coding region AF092	28	3e-20 7.1 <b>9.3</b>
Database: swissprot 79,449 sequences; 28,874,452 total letters		
Sequences producing significant alignments:	Score (bits)	E Value
sp P42087 HUTM_BACSU PUTATIVE HISTIDINE PERMEASE. sp P04775 CIN2_RAT SODIUM CHANNEL PROTEIN, BRAIN II ALPHA SUBU sp P42619 YQJF_ECOLI HYPOTHETICAL 17.2 KD PROTEIN IN EXUR-TDCC	26	7.1 9.2 9.2

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100104|lan|770RF104 Phage 77 ORF|34393-34551|1 (52 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score (bits)	_
gi 2315523 (AF016452) similar to the leucine-rich domains found gi 4377168 gb AAD18990  (AE001666) CT711 hypothetical protein [gi 3882171 dbj BAA34445  (AB018268) KIAA0725 protein [Homo sapi	29	5.4

Database: swissprot

79,449 sequences; 28,874,452 total letters

Sequences producing s	ignificant alignments:	Score (bits)	E Value
sp P04879 RRPP_VSVIG	RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48.		5.4
sp P04880 RRPP_VSVIM sp Q13946 CN7A HUMAN	RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48. HIGH-AFFINITY CAMP-SPECIFIC 3',5'-CYCLIC .		5.4 7.1
sp P35381 ATPA_DROME	ATP SYNTHASE ALPHA CHAIN, MITOCHONDRIAL P.		9.3
sp P54659 MVPB_DICDI sp P40397 YHXC BACSU	MAJOR VAULT PROTEIN BETA (MVP-BETA). HYPOTHETICAL OXIDOREDUCTASE IN APRE-COMK.		9.3 9.3

Score

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173

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BLASTP 2.0.8 [Jan-05-1999]
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Query= sid|122748|lan|770RF182 Phage 77 ORF|29268-29564|3 (98 letters)

Database: nr

393,678 sequences; 120,452,765 total letters

Sequences producing significant alignments:	(bits)	Value
gi 3341947 dbj BAA31913.1  (AB009866) orf 39 [bacteriophage phi.gi 1084792 pir  S54091 hypothetical protein YPR070w - yeast (Sa.gi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN.gi 744518 prf  2014422A FKBP-rapamycin-associated protein [Homo.gi 5051381 emb CAB44736.1  (AL049653) dJ647M16.2 (FK506 binding.gi 4826730 ref NP_004949.1 pFRAP1  FK506 binding protein 12-rap.gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa.gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa.gi 328239 (U88966) PRAPAMYCIN ASSOCIATED PROTEIN ASSOCIATED PROTEIN ASSOCIATED PROTEI	35 32 32 32 32	1.1 1.1 1.1
Database: swissprot 79,909 sequences; 29,054,478 total letters		
•		
•	Score	Æ
Sequences producing significant alignments:	Score (bits)	
Sequences producing significant alignments:	(bits)	
Sequences producing significant alignments:  sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP)	(bits)	Value
Sequences producing significant alignments:  sp!P42345 FRAP HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP)	(bits) 32 32	Value
Sequences producing significant alignments:  sp P42345 FRAP HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) sp P42346 FRAP RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R. sp P40557 YIA5_YEAST PUTATIVE DISULFIDE ISOMERASE YIL005W PREC. sp Q24118 LIO DROME LINOTTE PROTEIN.	(bits) 32 32 29 28	Value 0.29 0.29 3.3 4.4
Sequences producing significant alignments:  sp   P42345 FRAP HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) .  sp   P42346 FRAP RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R.  sp   P40557 YIA5 YEAST PUTATIVE DISULFIDE ISOMERASE YIL005W PREC.  sp   Q24118 LIO DROME LINOTTE PROTEIN.  sp   Q44363 TRAA AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	(bits) 32 32 29 28 28	Value 0.29 0.29 3.3 4.4 4.4
Sequences producing significant alignments:  sp P42345 FRAP HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) sp P42346 FRAP RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R. sp P40557 YIA5-YEAST PUTATIVE DISULFIDE ISOMERASE YIL005W PREC. sp Q24118 LIO_DROME LINOTTE PROTEIN. sp Q44363 TRAA AGRT6 CONJUGAL TRANSFER PROTEIN TRAA. sp P80034 ACH2 BOMMO ANTICHYMOTRYPSIN II (ACHY-II).	(bits)  32 32 29 28 28 28	Value 0.29 0.29 3.3 4.4 4.4
Sequences producing significant alignments:  sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) .  sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R.  sp P40557 YIA5_YEAST PUTATIVE DISULFIDE ISOMERASE YIL005W PREC.  sp Q24118 LIO_DROME LINOTTE PROTEIN.  sp Q44363 TRAA_AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	(bits)  32 32 29 28 28 28	Value 0.29 0.29 3.3 4.4 4.4

Table 6

1st position (5' end)	U	2nd p	osition A	G	3rd position (3' end)
	Phe	Ser	Tyr	Cys	U
n n	Phe	Ser	Tyr	Cys	С
U	Leu	Ser	Stop	Stop	Α
	Leu	Ser	Stop	Trp	G
	Leu	Pro	His	Arg	U
	Leu	Pro	His	Arg	С
$\mathbb{C}$	Leu	Pro	Gln	Arg	Α
	Leu	Pro_	Gln	Arg	G
	lle	Thr	Asn	Ser	U
۵	ile	Thr	Asn	Ser	С
A	lle	Thr	Lys	Arg	A
	Met	Thr	Lys	Arg	G
_	Val	Ala	Asp	Gly	U
<b>≈</b>	Val	Ala	Asp	Gly	С
G	Val	Ala	Glu	Gly	A
	Val	Ala	Glu	Gly	<u> </u>

#### Table 7

#### Bacteriophage 3A, complete genome sequence

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caaacgctag caacgcggat aaatttttca tgaaaggggg tctttatatg aagttaacaa aaaaacagct
        aaaagaatat atagaagatt acaaaaaatc tgatgacata ttaattaatt tgtatataga aacatatgaa
71
        ttttattgtc ggttaagaga tgaacttaaa aatagtgatt taatgataga gcatacaaac aaggctggtg
141
        cgagcaatat tattaagaat ccattaagca tagaactgac aaaaacagtt caaacactaa ataacttact
211
        caagtctatg ggtttaactg cagcacaaag aaaaaagata gttcaagaag aaggtggatt cggtgactat
281
        taaagtttta aatgaacctt caccaaaact attaacaaca tggtatgcag agcaagtcac tcaagggaaa
351
        ataaaaacaa gcaaatatgt tagaaaagaa tgtgagagac atcttagata tctagaaaat ggaggtaaat
421
       gggtatttga tgaagaatta gcgcatcgtc ctattcgatt tatagaaaag ttttgtaaac cttccaaagg
491
        atctaaacqt caacttgtat tacagccatg gcaacatttt attateggca gtttgtttgg ttgggttcat
561
        aaagaaacaa aactgcgcag gtttaaagaa gctttgatat ttatggggcg aaaaaatggt aaaaccaacca
631
        ctatttctgg ggttgctaac tatgctgtat cacaagatgg agaaaatggt gcagaaattc atttgttagc
701
        aaacgtaatg aaacaagcta ggattctatt tgatgaatct aaggcgatga ttaaagctag cccaaagctt
771
        gataaaaatt tcagaacatt aagagatgaa atccattatg acgcaacgat atcaaaaatt atgccccaag
841
        catcagatag cgataagtta gatggattga atacacacat ggggattttt gatgaaattc atgaatttaa
911
        agactataaa ttgatttcag ttataaaaaa ctcaagagct gcaaggttac aacctcttct catctacatt
981
1051
        acgacagcag ggtatcaatt agatggtcca cttgttgata tggtagaagc gggaagagac accttagatc
        aaatcataga agacgaaaga actttttatt atttagcatc tttggatgat gacgatgata ttaatgattc
1121
1191
        qtcqaactqq ataaaagcaa atcccaactt aggtgtctct ataaatttag atgagatgaa agaagagtgg
        gaaaaagcta agagaacacc agctgaacgt ggagatttta taaccaaaag gtttaatatc tttgctaata
1261
        atgacgagat gagttttatt gattacccaa cactccaaaa aaataatgaa attgtttctt tagaagagct
1331
1401
        ggaaggcaga ccgtgcacga ttggttatga tttatcagaa acagaggact ttacagccgc gtgtgctact
        tttgcgttag ataatggtaa agttgcagtt ttatcgcatt catggattcc taagcacaaa gttgaatatt
1471
1541
        ctaacqaaaa aatacctat aqaqaatggg aagaagatgg cttattaaca gtgcaagata agccttatat
       tgactaccaa gatgttttaa attggataat taagatgaat gagcattatg tagtagaaaa aattacttat
1611
1681
        gatagagcga acgcattcaa actaaatcaa gagttaaaaa attacgggtt tgaaacggaa gaaacaagac
        aaggagettt gaeettgage eetgeattga aggatttaaa agaaatgttt ttagatggga aaataatatt
1751
1821
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1891
        ttgccgtcta agcaaagcag atatcgtaaa atagatggct ttgcagcatt tttaaacaca tatacagata
1961
       ttatgaataa agttgtttct gatagtggtg aaggaaacat agagtttatt agtattaaag acataatgcg
       ttaaggaggt gaatgttatc gcaaaagaga atattgtcac acgcataaag aaaaaattga tagacaattg
2031
        gattgatcag tcaacttcta agctttatga ctttagccca tggaaaaata gatctttttg gggtgtaatt
2101
        aataatacgc ttgaaactaa tgaaacgata ttttcagcta ttacaaagtt atctaattcg atggctagtt
2171
       tgcccttgaa aatgtatgaa gattataaag tagttaatac agaagtatct gatttactta cagtgtcacc
2241
       gaataattot otgagoagtt ttgattttat taatcaaatt gaaacaatca gaaatgaaaa aggtaatgca
2311
       tatgtgctaa ttgaacgaga catctatcat caaccatcaa agcttttctt attaaatcca gatgttgttg
2381
       aaatgttaat tgaaaaccaa tcacgtgaac tttattattc cattcatgct gcaactggaa ataaattgat
2451
       tgttcataat atggacatgt tgcattttaa acacatcgtg gcatctaata tggtgcaagg cattagtccg
2521
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2591
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2661
       tttcaaacag tactatgaag aaaacggtgg aatattattc caagagcctg gtgttgaaat cgaaccgtta
2731
       cctaaaaaat atgtctctga agatatagtg gcaagcgaga atttaacaag agaaagagta gctaacgttt
2801
2871
       ttcaattgcc ctcagtattc ttaaatgcaa gatcaaatac aaatttcgcg aaaaatgaag agttaaacag
       attttacttg cagcatacct tattgccaat cgtcaaacag tatgaagaag aatttaatcg gaaactactt
2941
3011
       actaaaacaq acaqaqaaaa aaataggtat tttaaattta acgttaaatc ttatttaagg gctgatagtg
       caacacaagc agaagtgtac tttaaagcag ttcgtagtgg ttactacact ataaatgaca ttagagagtg
3081
       ggaagattta ccaccagttg aaggtggaga taagccgcta ataagcggtg atttataccc aattgacacg
3151
3221
       ccacttgaat taagaaaatc tttgaaaggt ggtgataaaa atgtcaatga aagctaagta ttttcaaatg
3291
       aaaagaaaat caaaaagtaa aggtgaaata tttatttatg gtgatattgt aagtgataaa tggtttgaaa
       gtgatgtaac tgctacagat ttcaaaaata aactagatga actaggagac atcagtgaaa tagatgttca
3361
       tataaattca totggaggca gtgtatttga agggcatgca atatacaata tgctaaaaat gcatcctgca
3431
       aaaattaata totatgtoga tgoottagog goatcaattg otagtgttat ogotatgagt ggtgacacta
3501
       tttttatgca caaaaatagt tttttaatga ttcataattc atgggttatg actgtaggta atgcagaaga
3571
       gttaagaaag acagcggatt tacttgaaaa aacagatgct gttagtaatt cagcttattt agataaagca
3641
       aaagatttag atcaagaaca cttaaaacag atgttagatg cagaaacttg gcttactgca gaagaagcct
3711
       tgtctttcgg cttgatagat gaaattttag gagctaatga aataactgct agtatctcta aagagcaata taagcgtttc gagaacgtcc cagaagattt aaagaaagat gtagacaaaa tcactaaaat cgatgatgta
3781
3851
3921
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       ttaaacgcga atgcgaaatt ttaaaaatga caatgagtta ttaggaggaa atgaaatgcc gacattatat
3991
4061
       qaattaaaac aatcettagg tatgattgga caacaattaa aaaataaaaa tgatgaattg agtcagaaag
       caacagaccc aaatattgat atggaagaca tcaaacaact agaaacagaa aaagcaggct tacaacaaag
4131
       atttaacatt gttgaaagac aagtaaaaga cattgaagaa aaagaaaaag cgaaagttaa agacacagga
4201
       gaagettate aatetttaaa tgateatgag aagatggtta aagetaagge agagttttat egteaegega oldsymbol{\bot}
4271
       ttttaccaaa tgaatttgaa aaacettcaa tggaggcaca acgtttatta cacgetttac caacaggtaa
4341
       tgattcaggt ggtgataagc tcttaccaaa aacactttct aaagaaattg tttcagaacc atttgctaaa
4411
       aaccaattac gtgaaaaagc tcgtctaact aacattaaag gtttagagat tccaagagtt tcatatactt
4481
       tagacgatga tgacttcatt acagatgtag aaacagcaaa agaattaaaa ttaaaaggtg atacagttaa
4551
       attcactact aataaattca aagtatttgc tgcaatttca gatactgtaa ttcatggatc agatgtagat
4621
```

ttagtaaact gggttgaaaa cgcactacaa tcaggtctag cagctaaaga acgtaaagat gccttagcag

\_\_\_\_

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ggtcaaagtt aaaaacgtat ggtgggaagt tagtcgagaa tattgttcaa gcaactgcaa gggatttact 33811 tgegatttet atagcaagge ttgaagcatt aggttttaaa atagttggee atgtecatga tgaagtaatt 33881 gtagaaatac ctagaggttc aaatggactt aaggaaatcg aaactatcat gaataagcct gttgattggg 33951 caaaaggatt gaatttgaat agtgacgggt ttacttctcc gttttatatg aaggattagg agtgtgattg 34021 catgcaacat caagcttata tcaatgcttc tgttgacatt agaattccta cagaagtcga aagtgttaat 34091 tacaatcaga ttgataaaga aaaagaaaat ttggcggact atttatttaa taatccaggt gaactattaa 34161 34231 aatataacgt tataaatatt aaggttttag atttagaggt ggaatgatgg ctagaagaaa agttataaga gtgcgtatca aaggaaaact aatgacattg agagaagttt cagaaaaata tcacatatct ccagaacttc 34301 ttagatatag atacaaacat aaaatgcgcg gcgatgaatt attgtgtgga agaaaagact caaaatctaa 34371 agatgaagtt gaatatatgc agagtcaaat aaaagatgaa gaaaaagaga gagaaaaaat cagaaaaaaa 34441 gcgattttga acctatacca acgaaatgtg agagcggaat atgaagaaga aagaaagaga agattgagac 34511 catggettta tgatggaacg ccacaaaaac attcacgtga tecgtactgg ttcgatgtca ettataacca 34581 aatgttcaag aaatggagtg aagcataatg agcgtaatca gtaacagaaa agtagatatg aacgaagcgc 34651 aagacaatgt taagcaacca gcgcactaca Catacggcga Cattgaaatt atagatttta tcgaacaggt 34721 tacggcacag tatccacctc aactagcatt cgcaataggt aatgcaataa aatacttgtc tagagcacct 34791 ttaaagaatg gtcatgagga tttagcaaag gcgaagttt acgtccaaag agcttttgac ttgtgggagt 34861 gatgaccatg acagatageg catgtaaaga atacttaaac caattttteg gatctaagag atatetgtat caggataacg aacgagtggc acatatccat gtagtgaatg gcacttatta ctttcacggg catatcgtac 35001 caggetggca aggegtgaaa aagacatttg atacagegga agagetegaa acatatataa agcaacatgg 35071 tttggaatac gaggaacaga agcaactaac tttattttaa ggagatagaa atgatgaaaa tcaaagttga 35141 aaaaataatg aaaatagacg aattaattaa gtgggcgcga gaaaatccgg agctatcatt tggcagaaaa 35211 tattatacaa cagacaaaaa tgatgaaaac tttatttact tcggtgtttt taaaaattgt tttaaaataa 35281 gcgattttat attagttaat gctactttta gtgtcaaagt tgaagaagaa gtaaccgaag aaactaagtt 35351 35421 tgataggttg tttgaagtgt acgagattca agaaggagtc tataaatctg catcatatga gaatgctagt ataaacgaac gtttaaaaaa tgacagaatt tttcttgcta aagcattcta catcttaaac gacgacctaa 35491 ctatgacgtt aatttggaaa gaaggagagt tgattaaata atggaacacg gttcaaaaga atattacgaa 35561 aagcaaagtg aatactggtt tgatgaagca agcaagtttt tgaagcaacg tgatgagctt attggagata tagctaagtt aagagagtgc aacaaagagc tggagaagaa agcaagtgca tgggataggt attgcaagag 35701 35771 cgttgaaaaa gatttaataa acgaatttgg caaagatggt gaaagagtta aatttggaat ggaattaaac aataaaattt ttatggagga agacgcaaat gaataaccgc gaacaatcg aacaatcagt tattagtgct 35841 agcgcgtata acggcaatga cacagaggga ttattaaaag agattgagga cgtgtataag aaagcgcaag 35911 35981 cgtttgatga aatacttgag ggtttaccta atgctatgca agatgcaatc aaagaagata ttggtcttga tgaagcagta ggaattatga cgggtcaagt tgtctataaa tatgaggagg agcaggaaaa tgactaacat 36051 attacaagtg aaactattat caaaagacgc tagaatgcca gaacgaaatc ataagacgga tgcaggttat 36121 gacatatttt cagctaaaac tgtcgtactt gagccacaag aaaaggcagt gatcaaaaca gatgtagctg taagcattcc agagggctat gtcggtttat taactagccg tagtggtgta agtagtaaaa cgcatttagt 36191 36261 gattgaaaca ggcaagatag acgcgggata tcatggtaat ttagggatta atatcaagaa tgataatgaa 36331 acgttagaga gtgaggatat gagtaacttt ggtcggagtc cttctggtat agatggaaaa tacaccctac 36401 tacctgtaac agataaattt ttatgtatga atggtagtta tgtcataaat aaaggcgaca aactagctca 36471 attggttate gtgcctatat ggacacctga actaaagcaa gtggaggaat tcgagagtgt ttcagaacgt 36541

36681 ggagcaaaag gcttcggaag tagcggagtg taaagacata ttagatcgag tcaaggaggt tttggggaag tgagtgacat gttagaaata tttttcatag ggtttggtgt ttatctattt tgtcgcatag gtattatttt totcaagagt aaaaagacta tacacacaaa cotatatgaa atgttgttga ttgctactat ctttgtgaca 36821 totacattig otgataaaca toaaaagacg catatottaa tagcattitt agtaatgitt titatgagta 36891 ageteaaaca ageteaaggg agetatgagg aatgacacaa tacetagtea caacatttaa agatteaaca 36961 ggacgtaagc atacacacat aactaaagct aagagcaatc aaaggtttac agttgttgat gcggagagta 37031 aagaagaagc gaaagagaag tacgaggcac aagttaaaag aaatgcagtt attaaattag ggcagttgtt 37101 tgaaaatata agggagtgtg ggaaatgact aaacaaatac taagattatt attettacta gcgatgtatg 37171 agctaggcaa gtatgtaact gagcaagtat atattatgat gacggctaat gatgatgcag aggcgccgag tgactttgaa aaaatcagag ctgaagttte atggtaatag ctattatcat ttttgaatta attatattaa 37241 37311 tgtgtttagc aatagcactg gaggtgttgt aaatatgtgg attgtcattt caattgtttt atctatattt 37381 ttattgatct tgttaagtag catttctcat aagatgaaaa ccatagaagc attggagtat atgaatgctt 37451 atcttttcaa 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42001	gatcaaagag	tatataaagc	tttacaaaat	aaagaactaa	cgcaagaaga	attgatgaaa	gctattaaag
42071	caagaatagc	taagcataag	taatggaggt	ataagatggg	aaaggcgtca	tatgatatta	agccaggaac
42141	atttaaatat	attgaatcag	aaatatataa	tttaaatgag	aacaagaaag	agataaatag	attgagaatg
42211	gagatactta	acccaacgaa	agaactagac	accaacattg	tgtatggacc	gttacaaaaa	ggagagccag
42281	ttagaacaac	tgagttaatg	gcgacaaggt	tattgactaa	taagatgtta	cgtaacttag	aagagatggt
42351	tgaagcagtt	gaaagtgagt	acttaaagtt	acctgaagat	cataagaaag	taataaggtt	aaagtattgg
42421	aataaagata	agaagctaaa	gatagaacaa	ataggggatg	cttgtcacat	gcatcgcaat	acagttacta
42491	caatacgaaa	gaactttgtt	aaagcgatag	cgtatcatgc	aggtatcaaa	taacattgtg	caaagattgt
42561	qcaaaaqqcc	tacaaatctg	tagtaatatg	atagtatcgg	aaagatgtat	aaagttatct	gaaagttata
42631	cqacataaat	acatgaggca	catcgctaag	cggtgtgtct	tttgttatgc	aatcaaagag	gtgtaagaga
42701	tgaccaagca	taataacatt	tataagcatg	gtcgtaagtc	atatcaatac	gattggttct	atcattcaaa
42771	agcatggaag	aagttaagag	agatagcatt	agatagagat	aattatcttt	gtcaaatgtg	tttacgcgaa
42841	gatattataa	cagatgcaaa	gattgtgcat	cacattattt	atgttgatga	agattttaac	aaagctttag
42911	acttagataa	tctaatgtca	gtttgttata	gctgtcataa	caaaattcat	gcaaatgata	atgacaaaag
42981	taatcttaag	aaaattagag	ttctaaaaat	ttaaataaaa	aaattattta	aataaaattt	tatgcccccc
43051	tgcccatcgg	cttaaaatgt	tttttcgccg	ggtaccggag	aggcc		

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Table 8

# Bacteriophage 3A ORFs list

SID	LAN	PRA	POS	a.a.	RBS sequence	STA	STO
100379	3AORF001	1	851513488	1657	acaggtacggatttaagaaaacttt	ttq	taa
100380	3AORF002	2	3766740114	815	tttaaaataatqaaaqqaqccqaac	atg	taa
100381	3AORF003	† <del>1</del>	3218834149	653	ttaaagaaattgaggtgtcaagaat	ttg	tag
100382	3AORF004	3	1745719370	637	gctattttattagaaaggaaggtgc	att	taa
	<del></del>	1	3342034	566	agaaaaaagatagttcaagaagaag	gtg	taa
100383	3AORF005	1	1557117154	527	cttttatttataggtaggtgattta	atg	taa
100384	3AORF006		1933720836	499	atgatagtaaaacaagttcagggcc	atg	taa
100385	3AORF007	2			_ · · · · · · · · · · · · · · · · · · ·	<del></del>	<del></del>
100386	3AORF008	3	2217623630	484	aatgatttagggtaggtgttgacca	atg	tga
100387	3AORF009	1	4072642093	455	gtaaatacttttataagaatggtag	gtg	taa
100388	3AORF010	] 3	1349114738	415	gaggcggactaacgctacagtaaaa	att	taa
100389	3AORF011	2	20393277	412	attaaagacataatgcgttaaggag	gtg	taa
100390	3AORF012	2	40015209	402	aaaaaagagaaaaaattaaacgcga	atg	taa
100391	3AORF013	1	3037931545	388	attttatgaatgcgagaataaatgc	atg	taa
100392	3AORF014	2	1473815562	274	attatatgggaggtttgactaatta	atg	tag
100393	3AORF015	3	32494034	261	cttgaattaagaaaatctttgaaag	gtg	tag
100394	3AORF016	-2	2558726273	228	aagaagctaagaaaaaaataaaaat	atg	tga
100395	3AORF017	3	67297370	213	ttaatttttaaggaggaaataagca	atg	taa
100396	3AORF018	3	2454025154	204	aataaaataaaaagtaggtgataag	atg	taa
100397	3AORF019	2	3156532128	187	ctataaaaattaaaaaggacggtat	ata	taa
100398	3AORF020	3	3615036713	187	gcagtaggaattatgacgggtcaag	ttg	taa
100399	3AORF021	2	2401124535	174	gtaataaaatttataaagaaaggaa	atg	tga
100400	3AORF022	-2	1242312938	171	taaagtaccagtagacaatgtaggt	att	tga
100401	3AORF023	1	74627917	151	aaaataaatcaaaggagaataattt	atg	taa
100402	3AORF024	1	2673127174	147	actaaataaaaataaggaggacact	atg	tga
100403	3AORF025	1	4210642543	145	taagcataagtaatggaggtataag	atg	taa
100404	3AORF026	2	3525535671	138	aagcaactaactttattttaaggag	ata	taa
	3AORF027	2	58886298	136	atattggctataatacagtggtttt	atc	taa
100405		-3	2784528255	136	ccttttaagatgtttatgatccttt	ctg	taa
100406	3AORF028	-	3434434748	134	ttaaggttttagatttagaggtgga	atq	taa
100407	3AORF029	3		131	tataaaaaaqqaqttggccagataa	atg	tag
100408	3AORF030	2	62996694			ata	taa
100409	3AORF031	1	2083321225	130	ttaacaaaattataggagtgagaaa	<del></del>	
100410	3AORF032	-2	3998440361	125	aaatagctgttagagggttacccct	ata	tag
100411	3AORF033	1	79578325	122	gaatatctgcgtcttttttatttga	ata	taa
100412	3AORF034	-2	2850628871	121	gttatcaacctaaggaggtgataac	atg	tag
100413	3AORF035	-2	1067111036	121	tcctagcttcctaacagcaccgcca	ata	tga
100414	3AORF036	2	3002030382	120	accaattttaaggaggagttaatca	atg	tga
100415	3AORF037	2	2181822165	115	aagtgtaagtaatagttaagagtca	gtg	tag
100416	3AORF038	-2	4200342347	114	gtactcactttcaactgcttcaacc	atc	tga
100417	3AORF039	2	2138621727	113	tccagaaaatctagagtcataggtt	ata	taa
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### Table 9

Bacteriophage 96, complete genome sequence

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42001	aggtaatagg	ataagtagtt	ttggtaagtt	tagcacgatt	tagtatttac	ttagaataaa	aattttgcta
42071	cattaattat	agggaatctt	acagttatta	aataactatt	tggatggatg	ttaatattcc	tatacacttt
42141	ttaacattac	tctcaagatt	taaatgtaga	taacaggcag	gtactacggt	acttgcctat	ttttttgtta
42211	taatgtaatt	acattaccag	taaccaatct	ggcttaaaac	cacatttccg	gtagccaatc	cggctatgca
42281	gaggacttac	ttgcgtaaag	tagtaagaag	ctgactgcat	atttaaacca	cccatactag	ttgctgggtg
42351	gttgttttt	atgttatatt	ataaatgatc	aaaccacacc	acctattaat	ttaggagtgt	ggttattttt
42421	tatgcaaaaa	aaacgaaaaa	aagttcataa	aaagtattgc	atatcacgtt	taaccgtgtt	ataataaggt
42491	ataccagttg	agaggaggat	aaaaagtgtt	agaaaatttt	aaaactatag	cagaaatcgc	cttttataca
42561	atgtcagcaa	ttgccatagc	gaaaacattg	aaaaaagacg	ataagtaagt	agacaagccc	gaaagggctg
42631	tctatatata	aattctaaca	ctaaaatact	atgaaaacaa	tttacattat	tttaatcatt	cttatttgga
42701	taaacgtgtt	tttaggcaac	gatataagta	aaagtgttgt	tgcactgctt	actactttac	tgcttatcaa
42771	tttatggaag	agggataaaa	atgacagcaa	taaaagaaat	aattgaatca	atagaaaagt	tattcgaaaa
42841	agaaacggga	tataaaattg	ctaaaaattc	cggattacca	tatcaaactg	tgcaagattt	aagaaatgga
42911	aaaacatctt	tatcagatgc	cagatttaga	acgataataa	agttatacga	gtatcaaaga	tcgcttgaaa
42981	acgaagaaga	taaataaaag	gagccaaaaa	tatgtttgtt	acaaaagaag	aatttaaaac	tttgaatgta
43051	aaagaagtat	ttgaatcagg	taaaaacttt	ataaaaatta	cagatggaag	acatgcaata	tattgggtaa
43121	atgatagata	cgtagtactt	gaccataaaa	aaggcgattt	gtacccgcaa	aaagcatacc	caaaatatat
43191	caaaagaaaa	ttagtaagtt	aaataattag	aaaaccacgt	cttaattgac	gtggttattt	tttaggtttg
43261	cgcgtgtcaa	atacgtgtca	atttagttct	atttctttag	ttttcttct	aaacttaatt	gcttgtaaac
43331	cgcatagtta	taggcttttc	agctatatac	caagataaga	tttatcccgc	cgtctccata	aaaatatgct
43401	tggaaacctt	gatttaatgg	ggttttaatc	tagcaagtgt	caaatatgtg	tcaagaaaat	aattttctga
43471	cacgttgacc	ttgctctttt	ttatgttcat	caagtaagtg	agagtaggtg	tctaaagtta	tagatatatt
43541	ataatggcct	aatcttttgc	taatatattc	aatagg			

Table 10

# Bacteriophage 96 ORFs list

SID	LAN	FRA	POS	a.a.	RBS sequence	STA	STO
100733	960RF001	1	2599929142	1047	ccttgaatcgaaaggaggttagcct	ttg	taa
100734	960RF002	1	3200833906	632	tttttacgactaaaggaggcaacca	atg	taa
100735	960RF003	1	3010931995	628	ttatattttagataaggagtagcct	atg	taa
100736	960RF004	1	3676038634	624	attttgattgaaatgaggtgcatac	atg	taa
100737	960RF005	3	3390335729	608	gtttattcgaaggaaaggtggttga	ata	taa
100738	960RF006	2	4058942043	484	aatgatttagggtaggtgttgacca	atg	tag
100739	960RF007	1	1865220091	479	tatacacacatactaaacctgaacg	att	tga
100740	960RF008	2	896010201	413	tggcagaatttgggggcgataacga	atg	tga
100741	960RF009	2	1744718670	407	gacgcaataacggaagtgatcgtca	atg	tga
100742	960RF010	1	3864739819	390	taaatataaataaagaggtgtgtaa	atg	tga
100743	960RF011	-1	1191195	358	gtagctcgcctacccttattatttt	ttg	tga
100744	960RF012	2	2004521013	322	tttaatgacaaattacctgacatag	atg	tga
100745	960RF013	3	2915730098	313	acttattataagggaggtttgttag	ttg	taa
100746	960RF014	1	2192522839	304	agaaaataaagtgaggtaataaaat	atg	tag
100747	960RF015	1	58126591	259	atacacggtaaaggtgggagaatag	atg	taa
100748	960RF016	1	78528607	251	aataaaatgttgaaaggagagaaaa	atg	taa
100749	960RF017	3	34444190	248	aaatttaacattaatatcactttaa	gtg	taa
100750	960RF018	-3	2828129000	239	taagctatgttgaacatcgctagtc	atg	tga
100751	960RF019	3	71887859	223	tttaccgttctaggacgtggtttaa	atg	taa
100752	960RF020	3	2132421908	194	gaagggcaaaaaggagttttgatat	atg	taa
100753	960RF021	3	66127175	187	attaaaaattaattaaaaggacggt	ata	tag
100754	960RF022	2	2453625093	185	aaagaaaaacgaaggagtgtattaa	atg	taa
100755	960RF023	1	52755811	178	catgaaatggtaggaggtatgaaaa	gtg	tag
100756	960RF024	3	1448115014	177	taaaacgataggagataacgaataa	atg	taa
100757	960RF025	2	2515725666	169	ataaaaaaattgaaaagaggtatat	att	taa
100758	960RF026	-3	1508415590	168	tcattcttaacatagcccttaattc	atg	tga
100759	960RF027	-1	12291732	167	aatagcaaataaaggagtgtaaaac	atg	taa
100760	960RF028	1	1696017454	164	aaggcgtgtgatacagtgaaaacaa	ttg	taa
100761	960RF029	-1	17362227	163	tatgagaaaaggagtcatataaaag	atg	taa
100762	960RF030	1	2553125995	154	ttttcaagagggagagtcgctcgta	ctg	tag
100763	960RF031	2	2363324097	154	tttagtattgaaggtgattctgtag	atc	tag
100764	960RF032	-2	22482706	152	ataagacaccaaaggggtttggcgc	atg	tga
100765	960RF033	-3	3914739605	152	agcatataaatcgtttagtgtttgt	ttg	taa
100766	960RF034	2	1318113615	144	tagaagtcgaaaaagtggaggcaat	ata	taa
100767	96ORF035	2	1062811053	141	gagctaggattgcaagcaacgatat	ttg	tga
100768	960RF036	2	2411024535	141	gtatttttcatagaggtggttaaat	atg	taa
100769	960RF037	1	1258312996	137	atgaggaacagaagcaaccaacttt	att	tga
100770	960RF038	1	1562816032	134	atgttaagaatgatgcctagtttaa	ttg	taa
100771	96ORF039	3	3981640220	134	ctaatacactttacttaattaaggg	gtg	taa
100772	960RF040	-3	2752827932	134	tttccataaataaacgaggacacca	atg	tga
100773	960RF041	3	1620616607	133	gatgagggggggggtgtcagagtag	atg	tga
100774	960RF042	2	3572036106	128	aagttactataactaaaattatggg	gtg	taa
100775	960RF043	-2	3571336081	122	ttaaacgtcccctcagtatttgtt	ttg	taa
100776	960RF044	-2	94609828	122	agtatccatcagttgaagataatct	ata	taa
100777	960RF045	-3	51395504	121	ttctttttgtattctgtaatattca	att	tga
100778	960RF046	2	1151311872	119	aagtaaatgtatagaggtggaataa	atg	taa
100779	960RF047	2	2299123350	119	gtcgtactacgtctgataagagcga	gtg	tag
100780	960RF048	3	86078963	118	tggaaaaagaattgagtgatgacta	atg	tga
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	<del></del>				<del>,</del>	<del>,</del>	
101082	960RF350	-2	4080140911	36	accattccaattttgcccatatgat	gtg	tag
101083	960RF351	-2	3895339063	36	tatcttttaaaattctcgtaatagc	atc	taa
101084	960RF352	-2	3158531695	36	tagetgtcatcactagtatttttga	atc	taa
101085	960RF353	-2	2455024660	36	atagtccgttttaccgcctcgtact	att	tag
101086	960RF354	- 2	2008320193	36	atcatcattttgatatttctcaaac	ata	tga
101087	960RF355	- 2	9911101	36	gcatcttggcagtacgacgtaaaac	atc	tag
101088	960RF356	-3	3814838258	36	taagaaagcgtgcgcgatcaaataa	att	tga
101089	960RF357	-3	87908900	36	tgaagttatctagcgctattttct	ttg	tag
101090	960RF358	-3	44584568	36	ttcataaaagtattctttgtagtat	atg	tag
101091	960RF359	1	46664773	35	ttatcaaaatatacaacttaattaa	atc	tag
101092	960RF360	1	1156911676	35	ataaatttaccgaacatgaaaatga	att	tga
101093	960RF361	2	61226229	35	ggaaaacaaattgatgttgtagtga	ttg	taa
101094	960RF362	-1	4041840525	35	ttcgtaggtgtcattacttctttaa	ttg	tag
101095	960RF363	-1	3435834465	35	gttttgcttgatttcgatttgttga	atg	tga
101096	960RF364	-1	2065420761	35	ctatttccactgattccccatctaa	atg	tga
101097	960RF365	-1	84238530	35	tctttttagagttacgaggtttca	att	tag
101098	960RF366	-1	24022509	35	tgacgtatggcaacattttagatca	atc	taa
101099	960RF367	-2	3660736714	35	aaaataaaagccagtgccgaagca	ctg	tag
101100	960RF368	-2	2706127168	35	caaatcgtcctgcagcgttcaataa	atc	tag
101101	960RF369	-2	2647026577	35	atgagttgttaagtttaccccaaat	atc	taa
101102	960RF370	-2	1032710434	35	ccgtgccatcttctcggtataagta	ata	taa
101103	960RF371	-2	86508757	35	gggtacgggttgttactgttgatat	atc	taa
101104	960RF372	-3	1438214489	35	gttcttttaattgatctactgttaa	att	taa
101105	960RF373	- 3	81518258	35	atgtttgttagtctctgtgtagtct	atg	taa
101106	960RF374	-3	50075114	35	aaacgatttaagtggaacattattc	ata	taa
101107	960RF375	2	3056330667	34	cgattagaaatctttaaaaaaggac	ttg	tga
101108	960RF376	-1	1991620020	34	tctatgtcaggtaatttgtcattaa	att	taa
101109	960RF377	-1	92369340	34	cttttctgttagtaattgtttttaa	atc	taa
101110	960RF378	-1	90269130	34	actctttatctttagttgcttttaa	ata	tag
101111	960RF379	-2	2844728551	34	cttttgtgataataaagtttagtgc	ttg	tga
101112	960RF380	-3	4032940433	34	ccatttaccttcttgagatgttgga	ttg	tga
101113	960RF381	-3	3980139905	34	caaaagatgaaggctttttccatac	ttg	taa
101114	960RF382	-3	3383133935	34	atgttgtttgtaactcgattaagtt	atc	tga
101115	960RF383	-3	3368733791	34	gttattacgtcttaatacttgtgtt	gtg	tag
101116	960RF384	-3	1353013634	34	tatacgcactagtactgatcactga	ttg	taa
101117	960RF385	-3	38433947	34	tttgattgattgttctagttaagaa	att	taa
101118	960RF386	1	1225612357	33	agtcataaagaagttagcaatgtga	ttg	tag
101119	960RF387	2	22072308	33	tccaagactctttaactgttaactt	atc	tag
101120	960RF388	2	25192620	33	attgttgaatttcgattgatctaaa	atg	tga
101121	960RF389	2	2251722618	33	agaagtaaaatgcgtaatgctttag	atg	tag
101122	960RF390	2	2730227403	33	ttccaaaattgggctaatagtgtag	ctg	taa
101123	960RF391	2	3238432485	33	actaaaaaggttgagaaagctgtag	atg	taa
101124	960RF392	2	3928739388	33	aaaaacggtactgtagtatcaatca	atc	tag
101125	960RF393	3	1815318254	33	gtagtatatgccgactttgatttga	atg	taa
101126	960RF394	3	2418924290	33	tcagaccctaacattaacaaactag	ttg	tga
101127	960RF395	-1	1526615367	33	tcgataatttgtatagcttgtttta	atg	tag
101128	960RF396	- 2	3223932340	33	ttttagtgaaagcatctagtgttga	ata	tag
101129	960RF397	- 2	1612316224	33	ttatgtgtgcctatcatattaacaa	ttg	tag
101130	960RF398	-2	1364813749	33	tctttaactgaatgttgaatagcat	ttg	tag
101131	960RF399	-2	1098711088	33	acttctgtaggtattcttatatcaa	ttg	tga
101132	960RF400	-2	33823483	33	cttactggtaattcttcaaaattaa	atg	taa
101133	960RF401	- 3	4079440895	33	ccatatgatgtgaaagtgtttaaat	ttg	taa
101134	960RF402	- 3	3997840079	33	atattcctaaatcacttgaacctaa	att	tga
101135	960RF403	-3	3860738708	33	atcttcagtgtaaaatcgacagcca	atg	tag
101136	960RF404	-3	2128821389	33	cagacaccgtcttaagtccctttag	ata	taa

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#### Table 11

#### SEQUENCE INFORMATION FOR PHAGES MATCHING WITH TABLE 1

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M32695
  Bacteriophage PM2 nuclease cleavage site
  gi|166145|gb|M32695|BM2NCS [166145]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M32693
  Bacteriophage PM2 Hind III fragment 4
  gij166144|gb|M32693|BM24HIND3 [166144]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
  Bacteriophage PM2 Hind III fragment 4
  gi|166144|gb|M32693|BM24HIND3 [166144]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M32694
  Bacteriophage PM2 Hind III fragment 3
  gi|166143|gb|M32694|BM23HIND3 [166143]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
M26134
  Bacteriophage PM2 structural protein gene containing purine/pyrimidine rich
  regions and anti-Z-DNA-IgG binding regions, complete cds
  gij289360|gb|M26134|BM2PROTTV [289360]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
J02452
  bacteriophage fi 3'-terminal region ma
  gi|215409|gb|J02452|PFITR3 [215409]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
AF020798
  Bacteriophage Chp1 genome DNA, complete sequence
  gi|217761|dbj|D00624|BCP1 [217761]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 12 protein links, or 1 genome link)
X72793
  Clostridium botulinum C phage BONT/C1, ANTP-139, ANTP-33, ANTP-17, ANTP-70
  genes and ORF-22
  gi|516171|emb|X72793|CBCBONT [516171]
  (View GenBank report, FASTA report, ASN, 1 report, Graphical view, 1 MEDLINE link, 6 protein links, or 4 nucleotide neighbors)
X51464
   Clostridium botulinum D Phage C3 gene for exoenzyme C3
   gi[14907|emb|X51464|CBDPE3 [14907]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
D90210
   Bacteriophage c-st (from C. botulinum) C1-tox gene for botulinum C1 neurotoxin
   gi|217780|dbj|D90210|CSTC1TOX [217780]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
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S49407
   type D neurotoxin [bacteriophage d-16 phi, host = C. botulinum, type D, CB16, Genomic, 4087 nt]
   gi|260238|gb|S49407|S49407 [260238]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
 X53370
   Bacteriophage phi29 temperature sensitive mutant TS2(98) DNA polymerase gene
   gij15733|emb|X53370|POTS298 [15733]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)
   Bacteriophage phi29 temperature sensitive mutant TS2(24) DNA polymerase gene
   gi|15731|emb|X53371|POTS224 [15731]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)
X05973
   Bacteriophage phi29 prohead RNA
  gi|15680|emb|X05973|POP29PRO [15680]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 4 nucleotide neighbors)
V01155
  Left end of bacteriophage phi-29 coding for 15 potential proteins Among
  these are the terminal protein and the proteins encoded by the genes 1, 2 (sus), 3, and (probably) 4
  gi|15659|emb|V01155|POP29B [15659]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 16 protein links, or 16 nucleotide neighbors)
  Bacteriophage phi-29 left origin of replication
  gi|312194|emb|X73097|BP29ORIL [312194]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
M14430
  Bacteriophage phi-29 gene-17 gene, complete cds
  gi|215321|gb|M14430|P29G17A [215321]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 6 protein links, or 8 nucleotide neighbors)
M14431
  Bacteriophage phi-29 gene-16 gene, complete cds
  gi|215319|gb|M14431|P29G16A [215319]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 7 nucleotide neighbors)
M20693
  Bacteriophage phi-29 DNA, 3' end
  gi|215343|gb|M20693|P29REPINB [215343]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
M21016
  Bacteriophage phi-29 DNA, 5' end
  gi|215342|gb|M21016|P29REPINA [215342]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
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M12456
   Bacteriophage phi-29 genes 9, 10 and 11 encoding p9 tail, incomplete, p10
   connector, complete, and pll lower collar, incomplete, respectively
   gi|215338|gb|M12456|P29P9 [215338]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)
 M14782
   Bacillus phage phi-29 head morphogenesis, major head protein, head fiber
   protein, tail protein, upper collar protein, lower collar protein, pre-neck-
   appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds
   gi|215323|gb|M14782|P29LATE2 [215323]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)
 M26968
   Bacteriophage phi-29 (from Bacillus subtilis) proteins p1 delta-1 genes, complete cds, and the sus1(629) mutation
   gi|341558|gb|M26968|P29P1D1A [341558]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
 J02448
   Bacteriophage fl, complete genome
   gi[166201]gb|J02448]F1CCG [166201]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, 205 nucleotide neighbors.
   or I genome link)
M24832
   Bacteriophage f2 coat protein gene, partial cds
   gil166228|gb|M24832|F2CRNACA [166228]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
J02451
  Bacteriophage fd, strain 478, complete genome
  gi|215394|gb|J02451|PFDCG [215394]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 MEDLINE links, 10 protein links, 204 nucleotide neighbors,
  or I genome link)
M34834
  Bacteriophage fr replicase gene, 5' end
  gi|166139|gb|M34834|BFRREGRA [166139]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors)
M38325
  Bacteriophage fr replicase gene, 5' end
  gi|166137|gb|M38325|BFRREGR [166137]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors)
M35063
  Bacteriophage fr coat protein replicase cistron (R region) RNA
  gi|166134|gb|M35063|BFRRCRRA [166134]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 3 nucleotide neighbors)
  alpha-atrial natriuretic factor/coat protein=fusion polypeptide [human.
  bacteriophage fr, expression vector pFAN15, PlasmidSyntheticRecombinant, 510 nt]
  gi|435742|gb|S66567|S66567 [435742]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 15 nucleotide neighbors)
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X15031

```
Bacteriophage fr RNA genome
   gi|15071|emb|X15031|LEBFRX [15071]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, 9 nucleotide neighbors,
   or I genome link)
 U51233
   Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable
   region light chain (IgM) mRNA, partial cds
   gi|1277150|gb|U51233|MMU51233 [1277150]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 1669 nucleotide neighbors)
U51232
   Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable region heavy chain (IgM) mRNA, partial cds
  gi|1277148|gb|U51232|MMU51232 [1277148]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 1073 nucleotide neighbors)
U02303
  Bacteriophage If1, complete genome
  gi|3676280|gb|U02303|B2U02303 [3676280]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, or 1 genome link)
V00604
  Phage M13 genome
  gi|14959|emb|V00604|INM13X [14959]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 205 nucleotide
neighbors)
A32252 ·
  Synthetic bacteriophage M13 protein III probe
  gi|1567340|emb|A32252|A32252 [1567340]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
A32251
  Synthetic bacteriophage M13 protein III probe
  gi|1567339|emb|A32251|A32251 [1567339]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
M12465
  Bacteriophage M13 mp10 mutations in lac operon
  gi|215210|gb|M12465|M13LACMUT [215210]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 215 nucleotide neighbors)
M24177
  Synthetic Bacteriophage M13 (clone M13.SV.B12) SV40 early promoter region DNA
  gi|209416|gb|M24177|SYNSVB12 [209416]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
  Synthetic Bacteriophage M13 (clone M13.SV.B11) SV40 early promoter region DNA
  gi|209415|gb|M24176|SYNSVB11 [209415]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
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## M24175 Synthetic Bacteriophage M13 (clone M13.SV.8) SV40 early promoter region DNA gi|208806|gb|M24175|SYNM13SV8 [208806] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 242 nucleotide neighbors) M19979 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207813|gb|M19979|SYN33M13M [207813] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 617 nucleotide neighbors) M19565 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207808|gb|M19565|SYN33M13H [207808] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 567 nucleotide neighbors) Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207807|gb|M19564|SYN33M13G [207807] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 12 nucleotide neighbors) M19563 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207806|gb|M19563|SYN33M13F [207806] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 262 nucleotide neighbors) Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207804|gb|M19561|SYN33M13D [207804] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 27 nucleotide neighbors) M19560 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207803|gb|M19560|SYN33M13C [207803] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link) M19559 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207802|gb|M19559|SYN33M13B [207802] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 227 nucleotide neighbors) M10568 Bacteriophage M13 replicative form II, replication origin, specific nick location gi|215220|gb|M10568|M13ORIB [215220] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 650 nucleotide neighbors) M10910 Bacteriophage M13 gene II regulatory region and M13sjl mutant gi|215209|gb|M10910|M13IREG [215209] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 72 nucleotide neighbors) M38295 Bacteriophage M13 HaeIII restriction fragment DNA gi|215208|gb|M38295|M13HAEIII [215208] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 67 nucleotide neighbors)

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.210
  E02067
    DNA encoding a part of Bacteriophage M13 tg 127
    gi|2170311|dbj|E02067|E02067 [2170311]
    (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
  J02467
    Bacteriophage MS2, complete genome
    gi|215232|gb|J02467|MS2CG [215232]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 8 MEDLINE links, 4 protein links, 20 nucleotide neighbors,
    or I genome link)
 AJ004950
    Bacteriophage P1 ban gene
    gi|3688226|emb|AJ011592|BP1011592 [3688226]
    (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 protein link)
    Bacteriophage P1 structural lytic transgivcosylase (orf47), pep44b (orf44b),
   pep44a (orf44a), and pep43 (orf43) genes, complete cds; and pep42 (orf42) gene, partial cds
   gi|2661099|gb|AF035607|AF035607 [2661099]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 1 nucleotide neighbor)
 AJ000741
   Bacteriophage P1 darA operon
   gi|2462938|emb|AJ000741|BPAJ7641 [2462938]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors
X01828
   Bacteriophage P1 recombinase gene cin
   gi|15133|emb|X01828|MYP1CIN [15133]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
X98146
   Bacteriophage P1 DNA sequence around the Op88 operator
   gi|1359513|emb|X98146|BP1OP880P [1359513]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)
S61175
  imml operon: icd=cell division repressor, ant1=antirepressor {promoters
  P51a, P51b} [bacteriophage P1, Genomic, 728 nt]
  gi|385908|gb|S61175|S61175 [385908]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
X87824
  Bacteriophage P1 gene 26
  gi|861164|emb|X87824|XXBP1G26 [861164]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)
X15638
  Phage P1 DNA for lytic replicon containing promoter P53 and two open reading frames
  gi|15735|emb|X15638|PP1LREP [15735]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 24 nucleotide neighbors
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### X17512 Bacteriophage P1 DNA for immunity region immI gi|15479|emb|X17512|P1IMMUNIY [15479] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 4 nucleotide neighbors) X16005 Bacteriophage P1 c1 gene for P1c1 repressor protein gi|15477|emb|X16005|P1C1 [15477] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors) X03453 Bacteriophage P1 cre gene for recombinase protein gi|15135|emb|X03453|MYP1CRE [15135] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors ? X06561 Bacteriophage P1 c1 gene 5'-region gi|15128|emb|X06561|MYP1C1 [15128] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 6 nucleotide neighbors) V01534 Bacteriophage P1 genome fragment (IS2 insertion spot). This regions contains four unidentified reading frames and is known as insertion hot spot for IS2 insertion sequences gi|15118|emb|V01534|MYOVP1 [15118] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors) X56951 Bacteriophage P1 gene10 gi|406728|emb|X56951|BPP1GP10 [406728] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor) K02380 Bacteriophage P1 replication region including repA, parA, and parB genes and incA, incB, and incC incompatibility determinants gi|215652|gb|K02380|PP1REP [215652] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 MEDLINE links, 4 protein links, or 8 nucleotide neighbors) X87674 Bacteriophage P1 lydA & lydB genes gi|974763|emb|X87674|BACP1LYD [974763] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors) X87673 Bacteriophage P1 gene 17 gi|974761|emb|X87673|BACP117 [974761] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor) M16618 Bacteriophage P1 c1 repressor binding sites gi|215600|gb|M16618|PP1C1 [215600] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

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SEG_PPICIN
    Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
     gi|215607|gb||SEG_PP1CIN [215607]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
  K03173
     Bacteriophage P1 C invertible element, right end, and cixR recombination site
     gi;215606|gb|K03173|PP1CIN2 [215606]
    (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
  215605
    Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
    gi|215605|lcl|X01828 [215605]
    (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
 M25470
    Bacteriophage P1 tail fiber protein gene, complete cds
    gi|341349|gb|M25470|PP1TFPR [341349]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)
 M34382
   Bacteriophage P1 sim region proteins, complete cds
   gi|215661|gb|M34382|PP1SIM [215661]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 M81956
   Bacteriophage P1 R protein (R) gene, complete eds
   gi|215658|gb|M81956|PP1RP [215658]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)
 M37080
   Bacteriophage P1 mini-P1 plasmid origin of replication
   gi|215657|gb|M37080|PP1REPOR [215657]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 46 nucleotide neighbors)
M27041
   Bacteriophage P1 ref gene, complete cds
  gi|215650|gb|M27041|PP1REF [215650]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
L01408
  Bacteriophage P1 partition protein (parB) gene, 3' end
  gi|215642|gb|L01408|PP1PARB [215642]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 41 nucleotide neighbors)
SEG_PPIPAR
  Bacteriophage miniplasmid P1 parA gene, 5' end
  gi|215639|gb||SEG_PP1PAR [215639]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 48 nucleotide neighbors)
M36425
  Bacteriophage miniplasmid P1 parB gene, 3' end
  gi|215638|gb|M36425|PP1PAR2 [215638]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
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213 M36424 Bacteriophage miniplasmid P1 parA gene, 5' end gij215637|gb|M36424|PP1PAR1 [215637] (View GenBank report, FASTA report, ASN.1 report, or Graphical view) M11129 Bacteriophage P1 miniplasmid origin of replication region gi|215632|gb|M11129|PP1ORIM [215632] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 43 nucleotide neighbors) M25414 Bacteriophage P1 c1 repressor binding site, operator 88 (Op88) gij215631|gb|M25414|PP1OP88A [215631] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors) M25413 Bacteriophage P1 c1 repressor binding site, operator 68 (Op68) gi|215630|gb|M25413|PP1OP68A [215630] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link) M25412 Bacteriophage P1 c1 repressor binding site, operator 21 (Op21) gi|215629|gb|M25412|PP1OP21A [215629] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor) M10510 Bacteriophage P1 recombination site loxR gi|215628|gb|M10510|PP1LOXR [215628] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor) M10287 Bacteriophage P1 loxP X loxP recombination site gi|215627|gb|M10287|PP1LOXPX [215627] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors) M10494 Bacteriophage P1 recombination site loxP gi|215626|gb|M10494|PP1LOXP [215626] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 134 nucleotide neighbors) Bacteriophage P1 recombination site loxL gi|215625|gb|M10511|PP1LOXL [215625] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor) M10512 Bacteriophage P1 recombination site loxB gi|215624|gb|M10512|PP1LOXB [215624] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link) M10145 Bacteriophage P1 genome fragment with recombination site loxP gij215623|gb|M10145|PP1CREX [215623] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 21 nucleotide neighbors)

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M13327

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Bacteriophage P1 Cin recombinase activated cross over site, junction IV, clone pSHI326
     gi|215622|gb|M13327|PP1CN26IV [215622]
     (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
     Bacteriophage P1 Cin recombinase activated cross over site, junction II, clone pSHI326
    gi|215621|gb|M13325|PP1CN26II [215621]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1401 nucleotide neighbors)
  M13323
    Bacteriophage P1 Cin recombinase activated cross over site, junction IV, clone pSHI325
    gi|215620|gb|M13323|PP1CN25IV [215620]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
  M13321
    Bacteriophage P1 Cin recombinase activated cross over site, junction II, clone pSHI325
    gi|215619|gb|M13321|PP1CN25II [215619]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1058 nucleotide neighbors)
 M13324
   Bacteriophage P1 Cin recombinase activated cross over site, junction I, clone pSHI326
   gi|215618|gb|M13324|PP1CIR26I [215618]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
 M13319
   Bacteriophage P1 Cin recombinase activated cross over site, right junction, clone pSHI327
   gi|215617|gb|M13319|PP1CIN27R [215617]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
 M13320
   Bacteriophage P1 Cin recombinase activated cross over site, junction I, clone pSHI325
   gi|215616|gb|M13320|PP1CIN251 [215616]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
M13318
   Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI324
   gi|215615|gb|M13318|PP1CIN24L [215615]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1370 nucleotide neighbors)
  Bacteriophage P1 Cin recombinase activated cross over site, right junction, clone pSHI323
  gi|215614|gb|M13317|PP1CIN23M [215614]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1055 nucleotide neighbors)
M13316
  Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI323
  gi|215613|gb|M13316|PP1CIN23L [215613]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
M13315
  Bacteriophage P1 Cin recombinase activated cross over site, right junction, clone pSHI322
  gi|215612|gb|M13315|PP1CIN22R [215612]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
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M13314
  Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI322
  gi|215611|gb|M13314|PP1CIN22L [215611]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1401 nucleotide neighbors)
M13313
  Bacteriophage P1 Cin recombinase activated cross over site, right junction, clone pSHI321
  gi|215610|gb|M13313|PP1CIN21R [215610]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
M13312
  Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI321
  gi|215609|gb|M13312|PP1CIN21L [215609]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1058 nucleotide neighbors)
M16568
  Bacteriophage P1 c4 repressor gene, complete cds
  gi|215603|gb|M16568|PP1C4 [215603]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
M13326
  Bacteriophage P1 Cin recombinase activated cross over site, junction III, clone pSHI326
  gi|215602|gb|M13326|PP1C26III [215602]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1192 nucleotide neighbors)
M13322
  Bacteriophage P1 Cin recombinase activated cross over site, junction III, clone pSHI325
 gi|215601|gb|M13322|PP1C25III [215601]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1231 nucleotide neighbors)
  Bacteriophage P1 modulator protein (bof) gene, complete cds
  gi[215598|gb|J05651|PP1BOFY1 [215598]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
M33224
  Bacteriophage P1 regulatory protein (bof) gene, complete cds
  gi|215596|gb|M33224|PP1BOFFO [215596]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
  E.coli/bacteriophage P1 loxR recombination site
  gi|146647|gb|M10288|ECOLOXR [146647]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
M10289
  E.coli/bacteriophage P1 loxL recombination site
  gij146646|gb|M10289|ECOLOXL [146646]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
M10290
  E.coli loxB site, which can recombine with bacteriophage P1 loxP site
  gi|146645|gb|M10290|ECOLOXB [146645]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
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M10287
    Bacteriophage P1 loxP X loxP recombination site
    gi|215627|gb|M10287|PP1LOXPX [215627]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
  M74046
    Bacteriophage P1 pacA and pacB genes, complete cds
    gi|215634|gb|M74046|PP1PACAB [215634]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 M95666
    Bacteriophage P1 gene 10, doc and phd genes, complete cds
    gi|463276|gb|M95666|PP1PHDDOC [463276]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 4 protein links, or 1 nucleotide neighbor)
 M25604
   Bacteriophage Q-beta mutated autonomously replicating sequence MDV1 RNA fragment
   gi|556359|gb|M25604|PQBARSMUT [556359]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)
 V00643
   first half of the phage Q-beta gene for coat protein
   gi|15088|emb|V00643|LEQBET [15088]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
 M25167
   Bacteriophage Q-beta RNA fragment recovered from replicase binding complex
   gi|556362|gb|M25167|PQBREPLICB [556362]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
M24876
   Bacteriophage Q-beta replicase RNA, 5' end
  gij556360|gb|M24876|PQBREPLICA [556360]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
M25444
  Synthetic bacteriophage Q-beta DNA
  gi|209118|gb|M25444|SYNPQBTERM [209118]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)
M25463
  Bacteriophage Q-beta self-replicating microvariant (+) RNA
  gi|532489|gb|M25463|PQBMVSRRNA [532489]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
M25014
  Bateriophage Q-beta RNA replicase gene, 5'end, and maturation protein gene, 3' end
  gi|294316|gb|M25014|PQBREPLC [294316]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
M25065
  Bacteriophage Q-beta RNA sequence with putative stem loop
  gi|294315|gb|M25065|PQBLOOP [294315]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
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M10265
      Bacteriophage Q-beta RNA molecule with the ability to replicate extracellularly
      gi|215726|gb|M10265|PQBRNA [215726]
      (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)
    M24815
      Bacteriophage Q-beta specified replicase subunit RNA,
      gi|215725|gb|M24815|PQBREPL [215725]
      (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
    M25461
      Bacteriophage Q-beta plus-strand RNA, 5' terminus
      gi|215724|gb|M25461|PQBPS5E [215724]
      (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
   M25462
      Bacteriophage Q-beta plus-strand RNA, 3' terminus
     gi|215723|gb|M25462|PQBPS3E [215723]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 8 nucleotide neighbors)
   M24871
     Bacteriophage Q-beta nanovariant WSIII RNA
     gi|215722|gb|M24871|PQBNVWSIC [215722]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
   M24870
     Bacteriophage Q-beta nanovariant WSII RNA
     gi|215721|gb|M24870|PQBNVWSIB [215721]
     (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
  M24869
     Bacteriophage Q-beta nanovariant WSI RNA
     gi|215720|gb|M24869|PQBNVWSIA [215720]
     (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
  M10495
     Coliphage Q-beta MDV-1(+) RNA
    gi|215719|gb|M10495|PQBMDV1A [215719]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 10 nucleotide neighbors)
· J02484
    bacteriophage queta coat protein cistron first half
    gi|215717|gb|J02484|PQBCP5 [215717]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
  M57754
    Bacteriophage Q-beta minus strand RNA, 5' terminus
    gi|215716|gb|M57754|PQBBMS5E [215716]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 8 nucleotide neighbors)
    Bacteriophage Q-beta 5'-terminal region of the minus strand
    gi|215715|gb|M24297|PQB5END [215715]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)
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M10695
                                                         218
   Bacteriophage Q-beta, MDV-I RNA
   gi|215714|gb|M10695|PQB1IR [215714]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 12 nucleotide neighbors)
 M24827
   Bacteriophage R17 A protein gene, 5' end
   gi|216078|gb|M24827|R17RNACIS [216078]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
 M24829
   Bacteriophage R17 coat protein gene, 5' end
   gi|216075|gb|M24829|R17CP5 [216075]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
J02488
   bacteriophage r17 rna synthetase initiation site
   gi|216080|gb|J02488|R17RNASYN [216080]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 MEDLINE links, 2 protein links, or 6 nucleotide neighbors)
J02487
  bacteriophage r17 coat protein initiation site
  gi|216073|gb|J02487|R17COATP [216073]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
J02486
  bacteriophage r17 a protein initiation site
  gi|216071|gb|J02486|R17APROT [216071]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
M24826
  Bacteriophage R17 coat protein RNA fragment
  gi|216077|gb|M24826|R17CPRAA [216077]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
M24296
  Bacteriophage R17 3'-terminal fragment A RNA
  gi|216070|gb|M24296|R173TFA [216070]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 9 nucleotide neighbors)
ITFN
  structure refinement for a 24-nucleotide rna hairpin, nmr, minimized average
  structure ribonucleic acid, hairpin, bacteriophage r17 mol_id: 1; molecule: r17c; chain: null; engineered: yes
  gi|1942336|pdb|1TFN| [1942336]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 structure link)
IRPEA
  rna (5'-d(gpgpgpapcpupgpapcpgpapupcpapcpgp cpapgpupcpupapu-3') (24-mer rna
 hairpin coat protein binding site for bacteriophage r17) (nmr, minimized average structure)
 gi|1421020|pdb|1RHT| [1421020]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 structure link)
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M14428
    Bacteriophage S13 circular DNA, complete genome
    gi|216089|gb|M14428|S13CG [216089]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 12 protein links, 26 nucleotide neighbors.
    or I genome link)
  J05393
    Bacteriophage T1 DNA N-6-adenine-methyltransferase (M.T1) gene, complete cds
    gi|166163|gb|J05393|BT1NAMTA [166163]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 L46845
    Bacteriophage T2 frd3, frd2 genes, complete cds
   gi|951387|gb|L46845|PT2FRD32G [951387]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 protein links, or 17 nucleotide neighbors)
 L43611
   Bacteriophage T2 fibritin (wac) gene, complete cds
   gij903869|gb|L43611|PT2WAC [903869]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)
 M24812
   Bacteriophage T2 secondary structure RNA sequence
   gi|215796|gb|M24812|PT2RNA [215796]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
M22342
   Bacteriphage T2 DNA-(adenine-N6)methyltransferase (dam) gene, complete cds
   gi|215792|gb|M22342|PT2DAM [215792]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
S57515
   orf 61.2 {intergenic region between 41 and 61} [bacteriophage T2, Genomic, 323 nt]
   gi|298524|gb|S57515|S57515 [298524]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
X05312
  Bacteriophage T2 gene 38 for receptor recognizing protein
  gi|15197|emb|X05312|MYT2G38 [15197]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
X04442
  Bacteriophage T2 gene 37 for receptor recognizing protein
  gi|15195|emb|X04442|MYT2G37 [15195]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
  Bacteriophage T2 gene 32 mRNA for single-stranded DNA binding protein
  gi|15192|emb|X12460|MYT2G32 [15192]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 14 nucleotide neighbors)
X57797
  Bacteriophage T2 gene for gp12
  gi|14875|emb|X56555|BT2GP12 [14875]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 2 nucleotide neighbors)
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# X01755 Bacteriophage T2 tail fiber gene 36 gi|15189|emb|X01755|MYT2F36 [15189] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor) M14784 Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis protein and DNA packaging proteins, complete cds gi|215810|gb|M14784|PT3RE [215810] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 10 nucleotide neighbors) SEG\_PT3RNAPOL Bacteriophage T3 RNA polymerase III gene, 5' end gi|710559|gb||SEG\_PT3RNAPOL [710559] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors) M22610 Bacteriophage T3 RNA polymerase III gene, 3' end gi|340722|gb|M22610|PT3RNAPOL2 [340722] (View GenBank report, FASTA report, ASN.1 report, or Graphical view) M22609 Bacteriophage T3 RNA polymerase III gene, 5' end gi|340721|gb|M22609|PT3RNAPOL1 [340721] (View GenBank report, FASTA report, ASN. 1 report, or Graphical view) X05031 Bacteriophage T3 gene region 1-2.5 with primary origin of replication gi|15719|emb|X05031|POT3ORI [15719] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 11 protein links, or 5 nucleotide neighbors) X03964 Bacteriophage T3 early control region pos. 308-810 from genome left end gi|15718|emb|X03964|POT3EP [15718] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 20 nucleotide neighbors) X17255 Bacteriophage T3 gene 1 to gene 11 gij15682|emb|X17255|POT3111G [15682] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 36 protein links, 17 nucleotide neighbors, or I genome link) X15840 Phage T3 gene 10 gi|15625|emb|X15840|PODT3G10 [15625] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors) X02981 Bacteriophage T3 gene 1 for RNA polymerase gi|15561|emb|X02981|PODOT3P [15561] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors) J02503 bacteriophage t3 5' end, terminally redundant sequence (trs) gi|215816|gb|J02503|PT3TRS1 [215816] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# SEG PT3TRS

bacteriophage t3 5' end, terminally redundant sequence (trs)
gi|215818|gb||SEG\_PT3TRS [215818]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

#### J02504

bacteriophage t3 3' end, terminally redundant sequence (trs) gi[215817]gb]J02504|PT3TRS2 [215817] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

H YPERLINK http://www.rs.noda.sut.ac.jp/~kunisawa h t t p://www.rs.noda.sut.ac.jp/~kunisawa Bacteriophage T4 genomic database compiled by Arisaka et al.

#### X95646

Bacteriophage T5 DNA for region 60.5%-71% of the T5 genome gi|2791557|emb|AJ001191|BTJ001191 [2791557] (View GenBank report,FASTA report,ASN.1 report,Graphical view,7 MEDLINE links, 12 protein links, or 6 nucleotide neighbors.)

#### X56847

Bacteriophage T5 genomic region encoding early genes D10-D15 gij15407|emb|X12930|MYT5D10 [15407] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 4 nucleotide neighbors)

#### AF039886

Bacteriophage T5 subclone T5.5.3r5.18r, single pass sequence, genomic survey sequence gi|2811154|gb|AF039886|AF039886 [2811154] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039885

Bacteriophage T5 subclone T5.40f,41f, single pass sequence, genomic survey sequence gi|2811153|gb|AF039885|AF039885 [2811153] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF039884

Bacteriophage T5 subclone T5.26.fr, single pass sequence, genomic survey sequence gi|2811152|gb|AF039884|AF039884 [2811152] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF039883

Bacteriophage T5 subclone 10-T5.5.7F, single pass sequence, genomic survey sequence gi|2811151|gb|AF039883|AF039883 [2811151] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF039882

Bacteriophage T5 subclone 41-T5.5.4BF, single pass sequence, genomic survey sequence gi|2811150|gb|AF039882|AF039882 [2811150] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF03988

Bacteriophage T5 subclone 39-T5.5.4aF, single pass sequence, genomic survey sequence gi|2811149|gb|AF039881|AF039881 [2811149] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

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# AF039880

Bacteriophage T5 subclone 19-T5.7.2r, single pass sequence, genomic survey sequence gi[2811148]gb]AF039880[AF039880 [2811148] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF039879

Bacteriophage T5 subclone 18-T5.7.2F, single pass sequence, genomic survey sequence gi|2811147|gb|AF039879|AF039879 [2811147]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039878

Bacteriophage T5 subclone 11-T5.5.7R, single pass sequence, genomic survey sequence gi[2811146]gb]AF039878[AF039878 [2811146] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 nucleotide neighbors)

#### AF039877

Bacteriophage T5 subclone T5.4FR, single pass sequence, genomic survey sequence gi|2811145|gb|AF039877|AF039877 [2811145] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF039876

Bacteriophage T5 subclone 22-T5.16R, single pass sequence, genomic survey sequence gi|2811144|gb|AF039876|AF039876 [2811144] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF039875

Bacteriophage T5 subclone 21-T5.16R, single pass sequence, genomic survey sequence gi|2811143|gb|AF039875|AF039875 [2811143]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039874

Bacteriophage T5 subclone 21-T5.16F, single pass sequence, genomic survey sequence gi|2811142|gb|AF039874|AF039874 [2811142] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF039873

Bacteriophage T5 subclone 09-T5.6F, single pass sequence, genomic survey sequence gi|28|1141|gb|AF039873|AF039873 [2811141] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF039872

Bacteriophage T5 subclone 09-T5.6R, single pass sequence, genomic survey sequence gi|2811140|gb|AF039872|AF039872 [2811140] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 nucleotide neighbors)

# AF039871

Bacteriophage T5 subclone 04-T5.26.R, single pass sequence, genomic survey sequence gi|2811139|gb|AF039871|AF039871 [2811139]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF039870

Bacteriophage T5 subclone 13-T5.42F, single pass sequence, genomic survey sequence gi|2811138|gb|AF039870|AF039870 [2811138] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

WO 00/32825

X69460

```
Bacteriophage T5 ltf gene for L-shaped tail fibers
    gi|15415|emb|X69460|MYT5LTF [15415]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 1 protein link, or 4 nucleotide neighbors)
    Bacteriophage T5 D15 gene for 5' exonuclease
   gi|15413|emb|X03402|MYT5EXOG [15413]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
 Z11972
   Bacteriophage T5 tRNA-Tyr, tRNA-Glu, tRNA-Trp, tRNA-Phe, tRNA-Cys and
   tRNA-Asn genes, and ORFs 91aa, 90aa, 42aa and 172aa
   gi|15795|emb|Z11972|T56TRNAG [15795]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors)
 X03898
   Bacteriophage T5 genes for tRNA-His, -Ser and -Leu
   gi|15786|emb|X03898|STT5RN1 [15786]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 2 MEDLINE links)
X04177
   Bacteriophage T5 gene for transfer RNA-Gln
   gi|15421|emb|X04177|MYT5TRNQ [15421]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
X03899
   Bacteriophage T5 genes for tRNA-Val, -Lys, -fMet, -Pro and -Ile3
  gi|15787|emb|X03899|STT5RN2 [15787]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
X03798
  Bacteriophage T5 gene for tRNA-Asp (GUC)
  gi|15472|emb|X03798|NCT5TRDG [15472]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
  Bacteriophage T5 tRNA gene cluster (27.8%-22.4%)
  gi|15420|emb|Y00364|MYT5TRN [15420]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
X03140
  Bacteriophage T5 DNA with rho-dependent transcription terminator (Hind III-P fragment)
  gi|15417|emb|X03140|MYT5RHO [15417]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
Z35070
  Bacteriophage T6 DNA
  gi|535228|emb|Z35074|MYEREGBT6 [535228]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
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Bacteriophage T6 DNA ligase gene, complete cds gi|215991|gb|M38465|PT6LIG55 [215991]

# 224 AF060870 Coliphage T6 small subunit distal tail fiber (gene 36) gene, partial cds; and large subunit distal tail fiber (gene 37) and tail fiber adhesin (gene 38) genes, complete eds gi|3676458|gb|AF052605|AF052605 [3676458] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 protein links, or 2 nucleotide neighbors) Z35072 Bacteriophage T6 DNA encoding ORF19.1 gene and g19 gene gi|535232|emb|Z35072|MYTAILT6 [535232] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links) X12488 Bacteriophage T6 gene 32 mRNA for single-stranded DNA binding protein gi|15843|emb|X12488|MYT6G32 [15843] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors) Z78095 Bacteriophage T6 DNA (1506 bp) gi|1488562|emb|Z78095|BPHZ78095 [1488562] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors) Bacteriophage T6 DNA for Ip5, Ip6 gij535215|emb|Z35079|MY57BT6 [535215] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor) X68725 E.coli bacteriophage T6 gene for beta-glucosyl-HMC-alpha-glucosyl-transferase gi|296439|emb|X68725|ECT6 [296439] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor) X69894 Bacteriophage T6 alt gene for ADP-Ribosyltransferase gi|15422|emb|X69894|MYT6ADP [15422] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor) Bacteriophage T6 frd3, frd2 genes, complete cds gi|951390|gb|L46846|PT6FRD32G [951390] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 protein links) M27738 Bacteriophage T6 translational repressor protein (regA), complete cds gij215993|gbjM27738|PT6REGA [215993] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 5 nucleotide neighbors)

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

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V01146
    Genome of bacteriophage T7
    gi|431187|emb|V01146|T7CG [431187]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 13 MEDLINE links, 60 protein links, 105 nucleotide
    neighbors, or 1 genome link)
  X60322
    Bacteriophage alpha3 genes A, B, K, C, D, E, J, F, G, H
    gi|14775|emb|X60322|BACALPHA [14775]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 10 protein links, 22 nucleotide neighbors,
    or I genome link)
 X13332
    Bacteriophage alpha3 DNA for origin of replication
    gi|15093|emb|X13332|MLA3ORPL [15093]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
   Bacteriophage alpha3 gene for protein A part., finger domain
   gi|15092|emb|X12611|MIA3AFIN [15092]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 6 nucleotide neighbors)
 X15721
   Bacteriophage alpha3 deletion mutation DNA for the origin region (-ori) of replication
   gi|14774|emb|X15721|BA3DMOR9 [14774]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
X15720
  Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication
  gi|14773|emb|X15720|BA3DMOR8 [14773]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
X15719
  Bacteriophage alpha3 insertion mutant DNA for the origin region (-ori) of replication
  gi|14772|emb|X15719|BA3DMOR7 [14772]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 10 nucleotide neighbors)
X15718
  Bacteriophage alpha3 deletion mutation DNA for origin region (-ori) of replication
  gi|14771|emb|X15718|BA3DMOR6 [14771]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
X15717
  Bacteriophage alpha3 deletion mutatnt DNA for origin region (-ori) of replication
  gi|14770|emb|X15717|BA3DMOR5 [14770]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 9 nucleotide neighbors)
X15716
  Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
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(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 10 nucleotide neighbors)

gi|14769|emb|X15716|BA3DMOR4 [14769]

X15715

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Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of of replication
    gi|14768|emb|X15715|BA3DMOR3 [14768]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
  X15714
    Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
    gi|14767|emb|X15714|BA3DMOR2 [14767]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
  X15713
    Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication
    gi|14766|emb|X15713|BA3DMOR1 [14766]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
 X62059
    Bacteriophage alpha3 origin of cDNA synthesis (oriGA)
   gi|14763|emb|X62059|AL3ORIGA [14763]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
 X62058
   Bacteriophage alpha3 origin of cDNA synthesis (oriAA)
   gi|14762|emb|X62058|AL3ORIAA [14762]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
   Bacteriophage alpha3 origin of DNA replication
   gi|166103|gb|J02444|AL3ORI [166103]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)
M25640
   Bacteriophage alpha-3 H protein gene, complete cds
   gi|166101|gb|M25640|AL3HP [166101]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)
M10631
  Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
  gi|166099|gb|M10631|AL3CSA [166099]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
X00774
  Bacteriophage alpha-3 gene J sequence
  gi|15431|emb|X00774|NCBA3J [15431]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)
M25640
  Bacteriophage alpha-3 H protein gene, complete cds
  gi|166101|gb|M25640|AL3HP [166101]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)
M10631
  Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
  gi|166099|gb|M10631|AL3CSA [166099]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
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PCT/IB99/02040

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227
 J02459
   Bacteriophage lambda, complete genome
   gi|215104|gb|J02459|LAMCG [215104]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 87 MEDLINE links, 67 protein links, 190 nucleotide
    neighbors, or I genome link)
 J02482
   Bacteriophage phi-X174, complete genome
   gi|216019|gb|J02482|PX1CG [216019]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 23 MEDLINE links, 11 protein links, 26 nucleotide neighbors,
   or I genome link)
 J02454
   Bacteriophage G4, complete genome
   gi|215415|gb|J02454|PG4CG [215415]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 6 MEDLINE links, 11 protein links, 20 nucleotide neighbors,
   or I genome link)
X60323
   Bacteriophage phiK complete genome
   gij1478118|emb|X60323|BPHIKCG [1478118]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, 18 nucleotide neighbors, or 1 genome link)
L42820
  Bacteriophage BF23 tail protein (hrs) gene, complete cds
  gi|1048680|gb|L42820|BBFHRS [1048680]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
X54455
  Bacteriophage BF23 gene 17 and gene 18
  gi|14797|emb|X54455|BF231718G [14797]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)
M37097
  Bacteriophage BF23 DNA, right end of terminal repetition
  gi|166115|gb|M37097|BBFRIGH [166115]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
M37096
  Bacteriophage BF23 DNA, left end of terminal repetition
  gi|166114|gb|M37096|BBFLEFT [166114]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M37095
  Bacteriophage BF23 A2-A3 gene, complete cds, and A1 gene, 5' end
  gi|166110|gb|M37095|BBFA2A3 [166110]
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(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor)

# AF056281

Bacteriophage BF23 clone bf23.mac5/6.1, genomic survey sequence gi|3090930|gb|AF056281|AF056281 [3090930] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056280

Bacteriophage BF23 clone bf23.mac3, genomic survey sequence gi|3090929|gb|AF056280|AF056280 [3090929] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056279

Bacteriophage BF23 clone bf23.mac18/21.34, genomic survey sequence gi|3090928|gb|AF056279|AF056279 [3090928] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056278

Bacteriophage BF23 clone bf23.mac16/19.33, genomic survey sequence gi{3090927|gb|AF056278|AF056278 [3090927] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056277

Bacteriophage BF23 clone bf23.mac16/19-33, genomic survey sequence gij3090926|gb|AF056277|AF056277 [3090926] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056276

Bacteriophage BF23 clone bf23.mac12/9-9, genomic survey sequence gi|3090925|gb|AF056276|AF056276 [3090925] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056275

Bacteriophage BF23 clone bf23.mac11/14-24, genomic survey sequence gi|3090924|gb|AF056275|AF056275 [3090924] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056274

Bacteriophage BF23 clone bf23.57r64r, genomic survey sequence gi|3090923|gb|AF056274|AF056274 [3090923] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 3 nucleotide neighbors)

# AF056273

Bacteriophage BF23 clone bf23.54fr, genomic survey sequence gi|3090922|gb|AF056273|AF056273 [3090922] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056272

Bacteriophage BF23 clone bf23.47fr.mac10/7, genomic survey sequence gi|3090921|gb|AF056272|AF056272 [3090921] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056271

Bacteriophage BF23 clone bf23.23.66r, genomic survey sequence gi|3090920|gb|AF056271|AF056271 [3090920] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF056270

Bacteriophage BF23 clone bf23.23.64f, genomic survey sequence gi|3090919|gb|AF056270|AF056270 [3090919] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056269

Bacteriophage BF23 clone bf23.23.60r, genomic survey sequence gi|3090918|gb|AF056269|AF056269 [3090918] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056268

Bacteriophage BF23 clone bf23.23.60f, genomic survey sequence gi|3090917|gb|AF056268|AF056268 [3090917] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 nucleotide neighbor)

# AF056267

Bacteriophage BF23 clone bf23.23.59r, genomic survey sequence gi|3090916|gb|AF056267|AF056267 [3090916] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056266

Bacteriophage BF23 clone bf23.23.59f, genomic survey sequence gi|3090915|gb|AF056266|AF056266 [3090915] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056265

Bacteriophage BF23 clone bf23.23.56r, genomic survey sequence gi[3090914]gb[AF056265[AF056265 [3090914] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056264

Bacteriophage BF23 clone bf23.23.56f, genomic survey sequence gi|3090913|gb|AF056264|AF056264 [3090913] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056263

Bacteriophage BF23 clone bf23.23.68f55r, genomic survey sequence gi|3090912|gb|AF056263|AF056263 [3090912] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056262

Bacteriophage BF23 clone bf23.23.43fr.66f, genomic survey sequence gi|3090911|gb|AF056262|AF056262 [3090911] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056261

Bacteriophage BF23 clone bf23.23.2fr, genomic survey sequence gi|3090910|gb|AF056261|AF056261 [3090910] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF056260

Bacteriophage BF23 clone bf23.23.55.f, genomic survey sequence gi|3090909|gb|AF056260|AF056260 [3090909] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF056259

Bacteriophage BF23 clone bf23.23.53.r, genomic survey sequence gi|3090908|gb|AF056259|AF056259 [3090908] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056258

Bacteriophage BF23 clone bf23.23.53.f, genomic survey sequence gi|3090907|gb|AF056258|AF056258 [3090907] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056257

Bacteriophage BF23 clone bf23.23.52.r, genomic survey sequence gi|3090906|gb|AF056257|AF056257 [3090906] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### 4 F056256

Bacteriophage BF23 clone bf23.23.52.f, genomic survey sequence gi|3090905|gb|AF056256|AF056256 [3090905] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056255

Bacteriophage BF23 clone bf23.23.49.r, genomic survey sequence gi|3090904|gb|AF056255|AF056255 [3090904] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### A F056254

Bacteriophage BF23 clone bf23.23.49.f, genomic survey sequence gi|3090903|gb|AF056254|AF056254 [3090903] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056253

Bacteriophage BF23 clone bf23.23.48.r, genomic survey sequence gi|3090902|gb|AF056253|AF056253 [3090902] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056252

Bacteriophage BF23 clone bf23.23.48.f, genomic survey sequence gi|3090901|gb|AF056252|AF056252 [3090901] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056251

Bacteriophage BF23 clone bf23.23.44.r, genomic survey sequence gi|3090900|gb|AF056251|AF056251 [3090900] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056250

Bacteriophage BF23 clone bf23.23.41.f, genomic survey sequence gi|3090899|gb|AF056250|AF056250 [3090899] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056249

Bacteriophage BF23 clone bf23.23.22.a.r, genomic survey sequence gi|3090898|gb|AF056249|AF056249 [3090898] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056248

Bacteriophage BF23 clone bf23.23.22.a.f, genomic survey sequence gi|3090897|gb|AF056248|AF056248 [3090897] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

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231
  AF056247
     Bacteriophage BF23 clone bf23.23.68.r, genomic survey sequence
     gi|3090896|gb|AF056247|AF056247 [3090896]
     (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
  Z50114
     Bacteriophage BF23 DNA for putative tail protein gene
     gi|2464952|emb|Z50114|BF23LATE [2464952]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 protein link)
  D12824
    Bacteriophage BF23 genes for minor tail protein gp24 and major tail protein gp25, complete cds
    gi|520578|dbj|D12824|BBF2TAIL [520578]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)
  Z34953
    Bacteriophage K3 ip9, ip7 and ip8 genes
    gi|535261|emb|Z34953|MYK3IP978 [535261]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
    Bacteriophage K3 DNA for Ip3 and Ip4
    gi|535229|emb|Z35075|MYEORF64K [535229]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 X05560
   Bacteriophage K3 gene 38 for receptor recognizing protein
   gi|15112|emb|X05560|MYK3G38 [15112]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
 X04747
   Bacteriophage K3 gene 37 for receptor recognizing protein
   gi|15110|emb|X04747|MYK3G37 [15110]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
X01754
  Bacteriophage K3 tail fiber gene 36
   gi|15108|emb|X01754|MYK3F36 [15108]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
M16812
  Bacteriophage K3 't' lysis gene, complete cds
  gi|215503|gb|M16812|PK3LYST [215503]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
L46833
  Bacteriophage K3 frd3, frd2 genes, complete cds
  gi|951377|gb|L46833|PK3FRD32G [951377]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)
L43613
  Bacteriophage K3 fibritin (wac) gene, complete cds
```

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)

gi|903861|gb|L43613|PK3WAC [903861]

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X01753
     Bacteriophage Ox2 tail fiber gene 36
     gi|15122|emb|X01753|MYOX2F36 [15122]
     (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
   L43612
     Bacteriophage Ox2 fibritin (wac) gene, complete cds
     gi|903848|gb|L43612|OX2WAC [903848]
     (View GenBank report, FASTA report, ASN 1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)
   Z46880
     Bacteriophage OX2 stp gene
     gi|599663|emb|Z46880|BPOX2STP [599663]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
  X05675
    Bacteriophage Ox2 gene 38 for receptor-recognizing protein and flanking regions
    gi|15124|emb|X05675|MYOX2G38 [15124]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
  M33533
    Bacteriophage RB18 translational repressor protein (regA) and Orf43.1, complete cds
    gi|216083|gb|M33533|RB18REGA [216083]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
 AF033329
    Bacteriophage RB18 single-stranded binding protein (gene 32) gene, partial cds, and 5' region
   gi|2645788|gb|AF033329|AF033329 [2645788]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 11 nucleotide neighbors)
 M86231
   Bacteriophage RB69 gene 62, 3'end; RegA (regA) gene, complete cds
   gi|215354|gb|M86231|P6962REGA [215354]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
 AF033332
   Bacteriophage RB69 single-stranded binding protein (gene 32) gene, partial cds, and 5' region
   gi|2645794|gb|AF033332|AF033332 [2645794]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 12 nucleotide neighbors)
U34036
  Bacteriophage RB69 DNA polymerase (43) gene, complete eds
  gi|1237125|gb|U34036|BRU34036 [1237125]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
V01145
  Bacteriophage H1 genome fragment Each Thymine given in this sequence represents a HMU-residue
  (HMU = 5-hydroxymethyluracil)
  gi|15557|emb|V01145|PODOH1 [15557]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
X05676
  Bacteriophage M1 gene 38 for receptor recognizing protein and flanking regions
  gi|15114|emb|X05676|MYM1G38 [15114]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
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# AF034575 Bacteriophage M1 putative integrase (int) gene, complete cds, and attP region, complete sequence gi|2662472|gb|AF034575|AF034575 [2662472] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link) AF033321 Bacteriophage M1 single-stranded binding protein (gene 32) gene, partial cds, and 5' region gi|2645772|gb|AF033321|AF033321 [2645772] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors) Bacteriophage TuIa 37 and 38 genes for receptor-recognizing proteins 37 and 38 (respectively), partial cds gi|14860|emb|X55190|BPTUIA [14860] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors) AF033334 Bacteriophage Tulb single-stranded binding protein (gene 32) gene, partial cds, and 5' region gi|2645798|gb|AF033334|AF033334 [2645798] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 5 nucleotide neighbors) X55191 Bacteriophage TuIb 37 gene for receptor-recognizing protein 37 (partial cds), 38 gene for receptor-recognizing protein 38, and t gene (partial cds) gi|14863|emb|X55191|BPTUIB [14863] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors) X13065 Bacteriophage phi80 early region gil14800|emb|X13065|BP80ER [14800] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 8 protein links, or 6 nucleotide neighbors) D00360 Bacteriophage phi80 cor gene gi|217782|dbj|D00360|P8080COR [217782] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 protein link) X01639 Bacteriophage phi 80 DNA-fragment with replication origin gi|15828|emb|X01639|XXPHI80 [15828] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 25 nucleotide neighbors) X04051 Lambdoid bacteriophage phi 80 int-xis region (integrase-excisionase region) gi|15770|emb|X04051|STPH180X [15770] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor) X06751 Phage Phi80 DNA for major coat protein gi|15768|emb|X06751|STPHI80C [15768] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 11 nucleotide neighbors) X75949 Bacteriophage phi80 DNA for ORF x171.8 and ORF x171.28 gi|458811|emb|X75949|ECORF171B [458811] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 28 nucleotide neighbors)

L40418

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# Bacteriophage phi-80 gene, complete eds gi|1019107|gb|L40418|P80A [1019107] (View GenBank report, FASTA report, ASN 1 report, Graphical view, 1 MEDLINE link, or 1 protein link) M24831 Bacteriophage phi-80 Tyr-tRNA gene, 3' end gi|215363|gb|M24831|P80TGY [215363] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 43 nucleotide neighbors) M10670 Bacteriophage phi-80 replication origin gi|215361|gb|M10670|P80ORI [215361] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor) M24825 Bacteriophage phi-80 RNA fragment gi|215360|gb|M24825|P80M3A [215360] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor) M11919 Bacteriophage phi-80 cI immunity region encoding the N gene gi|215358|gb|M11919|P80CI [215358] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors) M10891 Bacteriophage phi-80 attP site DNA gi|215357|gb|M10891|P80ATTP [215357] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor) M19473 Bacteriophage 933J (from E.coli) proviral Shiga-like toxin type 1 subunits A and B genes, complete cds gi|215072|gb|M19473|J93SLTI [215072] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 2 protein links, or 20 nucleotide neighbors) Y10775 Bacteriophage 933W ileX, stx2A and stx2B genes gi|1938206|emb|Y10775|BP933ILEX [1938206] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 36 nucleotide neighbors) X83722 Bacteriophage 933W slt-IIB gene gi|1490229|emb|X83722|B933WSLT [1490229] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 20 nucleotide neighbors) X07865 Bacteriophage 933W slt-II gene for Shiga-like toxin typeII subunit A and B gi|14892|emb|X07865|BWSLTII [14892] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 protein links, or 29 nucleotide neighbors) M16625 Bacteriophage H19B (from E.coli) sltIA and sltIB genes encoding Shiga-like toxin I subunits A and B, complete cds gi|215043|gb|M16625|H19BSLT [215043] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 24 nucleotide neighbors)

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#### 1102466

gi|407285|gb|U02466|BHU02466 [407285] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor)

# M26291

Bacteriophage D108 regulatory DNA-binding protein (ner) gene, complete cds gi|166194|gb|M26291|D18NER [166194] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

#### M11272

Bacteriophage D108 left-end DNA gi|166193|gb|M11272|D18LEDNA [166193] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)

# M18902

Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete cds gi|166191|gb|M18902|D18KIL [166191] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

# M10191 Bacteriophage D108, left end with Mu A protein binding sites L1 and L2 gi|166190|gb|M10191|D18BSL [166190] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors) J02447 bacteriophage d108 gene a 5' end gi|166189|gb|J02447|D18AAA [166189] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link) V00865 Bacteriophage D108 fragment from genes A and ner (C-terminus of ner and N-terminus of A) gi|15437|emb|V00865|NCD108 [15437] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links) X01914 Bacteriophage IKe gene for DNA binding protein gi|14957|emb|X01914|INIKEDBP [14957] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors) AF064539 Bacteriophage N15, complete genome gi|3192683|gb|AF064539|AF064539 [3192683] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, 60 protein links, 26 nucleotide neighbors. or I genome link) U02303 Bacteriophage If1, complete genome gi|3676280|gb|U02303|B2U02303 [3676280] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, or 1 genome link) AF007792 Bacteriophage Mu late morphogenetic region gi|3551775|gb|AF007792|AF007792 [3551775] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor) U24159 Bacteriophage HP1 strain HP1c1, complete genome gi|1046235|gb|U24159|BHU24159 [1046235] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 6 MEDLINE links, 41 protein links, 8 nucleotide neighbors. or I genome link) Z71579 Bacteriophage S2 type A 5.6 kb DNA fragment gil1679806|emb|Z71579|BPHS1ADNA [1679806] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 MEDLINE links, 9 protein links, or 9 nucleotide neighbors) X53238 Klebsiella sp. bacteriophage K11 gene 1 for RNA polymerase gi|14984|emb|X53238|KSK11RPO [14984]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

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X85010
     Bacteriophage A511 ply511 gene
     gi|853748|emb|X85010|BPA511PLY [853748]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
   U29728
    Bacteriophage N4 single-stranded DNA-binding protein (N4SSB) gene, complete cds
    gi|939708|gb|U29728|BNU29728 [939708]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 1 protein link)
  J02445
    bacteriophage bol 3'-terminal region ma
    gi|166152|gb|J02445|BO1TR3 [166152]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleonde neighbors)
    Bacteriophage L5 (from Leuconostoc oenos) genome
    gi|289353|gb|L06183|BL5GENM [289353]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 genome link)
 AF074945
   Mycoplasma arthritidis bacteriophage MAVI, complete genome
   gij3511243|gb|AF074945|AF074945 [3511243]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 15 protein links, 3 nucleotide neighbors, or 1 genome link)
 L13696
   Bacteriophage L2 (from Mycoplasma), complete genome
   gi|289338|gb|L13696|BL2CG [289338]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 MEDLINE links, 14 protein links, or 1 genome link)
 X80191
   Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins
   gi|517237|emb|X80191|BPP7PR [517237]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 genome link)
M19377
  Bacteriophage Pf3 from Pseudomonas aeruginosa (New York strain), complete genome
   gi|215380|gb|M19377|PF3COMNY [215380]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 5 nucleotide neighbors)
M11912
  Bacteriophage Pf3 from Pseudomonas aeruginosa (Nijinegen strain), complete genome
  gi|215371|gb|M11912|PF3COMN [215371]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 9 protein links, 5 nucleotide neighbors, or 1
  genome link)
V00605
  Bacteriophage Pf1 gene encoding DNA binding protein
  gi|14970|emb|V00605|INOPF1 [14970]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 proteine link, or 1 nucleotide neighbor)
L05626
 Bacteriophage PR4 capsid protein (P6) gene, complete cds
 gi|215735|gb|L05626|PR4P6MAJA [215735]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
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#### D13409

Bacteriophage phiCTX (isolated from Pseudomonas aeruginosa) cosR, attP, int genes gi|217776|dbj|D13409|BPHCOSR [217776]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

#### D13408

Bacteriophage phiCTX (isolated from Pseudomonas aeruginosa) cosL, ctx genes gi|217775|dbj|D13408|BPHCOSLCTX [217775]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 3 nucleotide neighbors)

#### M24832

Bacteriophage f2 coat protein gene, partial cds gi|166228|gb|M24832|F2CRNACA [166228]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

#### S72011

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi[2618967]gb[AF017629]AF017629 [2618967]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

# AF017628

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gil2618964|gb|AF017628|AF017628 [2618964]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

#### AF017627

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618961|gb|AF017627|AF017627 [2618961]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

# AF017626

Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds gi|2618958|gb|AF017626|AF017626 [2618958]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

# AF017625

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618955|gb|AF017625|AF017625 [2618955]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

#### AF017624

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int)genes, partial cds gi[2618952]gb]AF017624[AF017624 [2618952]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

# AF017623

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618949|gb|AF017623|AF017623 [2618949]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

# AF017622

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618946|gb|AF017622|AF017622 [2618946]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017621 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618943|gb|AF017621|AF017621 [2618943] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) D26449 Bacteriophage PS17 FI gene for tail sheath protein (gpFI) and FII gene for tail tube protein (gpFII), complete cds gi|452162|dbj|D26449|BPSFIFII [452162] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 2 protein links) X87627 Bacteriophage D3112 A and B genes gi|974768|emb|X87627|BPD3112AB [974768] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 1 nucleotide neighbor) U32623 Bacteriophage D3 transcriptional activator CII (cII) gene, complete cds gi|984852|gb|U32623|BDU32623 [984852] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 1 nucleotide neighbor) L34781 Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and peptidoglycan hydrolase (lytA) gene, partial cds gi|511838|gb|L34781|BPHHOLIN [511838] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors) L14810 Bacteriophage P22 (gp10) gene, complete cds, and (gp26) gene, complete cds gi|294053|gb|L14810|P22GP1026X [294053] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors) Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators gi|1143407|emb|X87420|BPES18GEN [1143407] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 5 protein links, or 9 nucleotide neighbors) L42820 Bacteriophage BF23 tail protein (hrs) gene, complete cds gi|1048680|gb|L42820|BBFHRS [1048680] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 1 protein link, or 1 nucleotide neighbor) X14980 Bacteriophage PRD1 XV gene for protein P15 (lytic enzyme) gi|15802|emb|X14980|TEPRD1XV [15802] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 1 protein link, or 4 nucleotide neighbors) X06321 Bacteriophage PRD1 gene 8 for DNA terminal protein gi|15800|emb|X06321|TEPRD18 [15800] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 10 nucleotide neighbors) X14336 Filamentous Bacteriophage I2-2 genome gi|14920|emb|X14336|INBI22 [14920] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 9 protein links, 1 nucleotide neighbor, or 1 genome link }

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  L05001
    Bacteriophage X glucosyl transferase gene, complete eds
    gi|216044|gb|L05001|PXFCLUSYLT [216044]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
  M29479
    Bacteriophage p4 sid and psu genes partial cds, and delta gene, complete cds gij215701
    gb|M29479|PP4SDP [215701]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 protein links, or 4 nucleotide neighbors)
  SEG PP4PSUSID
    Bacteriophage P4 capsid size determination protein (sid) gene, 5' end
    gi|215698|gb||SEG_PP4PSUSID [215698]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
 M29650
   Bacteriophage P4 polarity suppression protein (psu) gene, complete cds
    gij215697|gb|M29650|PP4PSUSID2 [215697]
   (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
   Bacteriophage P4 capsid size determination protein (sid) gene, 5' end
   gi|215696|gb|M29651|PP4PSUSID1 [215696]
   (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
 M27748
   Bacteriophage P4 gop, beta, and cII genes, complete cds and int gene, 3' end
   gi|215691|gb|M27748|PP4GOPBC [215691]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 nucleotide neighbor)
K02750
   Bacteriophage IKe, complete genome
   gi|215061|gb|K02750|IKECG [215061]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 10 protein links, 4 nucleotide neighbors, or 1
   genome link )
L40418
  Bacteriophage phi-80 gene, complete cds
  gi|1019107|gb|L40418|P80A [1019107]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
AF032122
  Bacteriophage SfII integrase (int) gene, partial cds; and bactoprenol glucosyl transferase (bgt), and glucosyl transferase II (gttl)
  genes, complete cds
  gi|2465412|gb|AF021347|AF021347 [2465412]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 4 protein links, or 2 nucleotide neighbors)
M35825
  Bacteriophage SF6 fragment D lysozyme gene, complete cds
  gi|216105|gb|M35825|SF6LYZ [216105]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)
Z35479
  Bacteriophage C16 ip1 gene
  gi|534936|emb|Z35479|BC16IP1 [534936]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
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#### X12638

Bacteriophage 21 DNA for gene 2

gi|296141|emb|X12638|B21GENE2 [296141]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

#### X02501

Bacteriophage 21 DNA for left end sequence with genes 1 and 2

gi|15825|emb|X02501|XXPHA21 [15825]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

#### M65239

Bacteriophage 21 lysis genes S, R, and Rz, complete cds

gi|215466|gb|M65239|PH2LYSGEN [215466]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

#### M58702

Bacteriophage 21 late gene regulatory region

gi|215465|gb|M58702|PH2LATEGE [215465]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)

#### M81255

Bacteriophage 21 head gene operon

gi|215454|gb|M81255|PH2HEADTL [215454]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 10 protein links, or 4 nucleotide neighbors)

#### M23775

Bacteriophage 21 glycoprotein 1 gene, complete cds, and glycoprotein gene, 5' end

gij215451[gb]M23775[PH2GPA [215451]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

# M61865

Bacteriophage 21 excisionase (xis), integrase (int) and isocitrate dehydrogenase (icd), complete cds

gi|215448|gb|M61865|PH22XISAA [215448]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 9 nucleotide neighbors)

# S72011

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618967|gb|AF017629|AF017629 [2618967]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

#### AF017628

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gij2618964|gb|AF017628|AF017628 [2618964]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

# AF017627

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618961|gb|AF017627|AF017627 [2618961]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

# AF017626

Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds

gi|2618958|gb|AF017626|AF017626 [2618958]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

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# AF017625

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Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618955|gb|AF017625|AF017625 [2618955]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors

# AF017624

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618952|gb|AF017624|AF017624 [2618952]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

# AF017623

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618949|gb|AF017623|AF017623 [2618949]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

#### AF017622

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618946|gb|AF017622|AF017622 [2618946]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

# AF017621

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618943|gb|AF017621|AF017621 [2618943]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

# M57455

Bacteriophage 42D (clone pDB17) (from Staphylococcus aureus) staphylokinase gene, complete cds gi|215344|gb|M57455|P42STK [215344]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors)

#### Y12633

Bacteriophage 85 DNA, promoter sequence of unknown gene gi|2058285|emb|Y12633|B85PROM [2058285] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# X98146

Bacteriophage P1 DNA sequence around the Op88 operator gi|1359513|emb|X98146|BP1OP88OP [1359513]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

#### Y07739

Staphylococcus phage Twort holTW, plyTW genes gi|2764979|emb|Y07739|BPTWGHOLG [2764979]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 protein links)

#### L07580

Bacteriophage phi-11 rinA and rin B genes, required for the activation of Staphylococcal phage phi-11 int expression gi|166160|gb|L07580|BPHRINAB [166160]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

# M34832

Bacteriophage phi-11 integrase (int) and excisionase (xis) genes, complete cds gi|166157|gb|M34832|BPHINTXIS [166157]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

M20394

```
Bacteriophage phi-11 S.aureus attachment site (attP)
     gi|166156|gb|M20394|BPHATTP [166156]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
   X23128
     Bacteriophage phi-13 integrase gene
     gi|758228|emb|X82312|PHI13INT [758228]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 3 nucleotide neighbors)
  X61719
    S.aureus phi-13 lysogen right chromosome/bacteriophage DNA junction
    gi|46625|emb|X61719|SAP13RINC [46625]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
    S.aureus phi-13 lysogen left chromosomal/bacteriophage DNA junction
    gi|46624|emb|X61718|SAP13LJNC [46624]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
  X61717
    Bacteriophage phi-13 core sequence for attachment
    gi|14799|emb|X61717|BP13ATTP [14799]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 3 nucleotide neighbors)
 U01875
   Bacteriophage phi-13 putative regulatatory region and integrase (int) gene, partial cds
   gi|437118|gb|U01875|U01875 [437118]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, or 4 nucleotide neighbors)
X67739
   S.aureus Bacteriophage phi-42 attP gene
   gi|14809|emb|X67739|BPATTPA [14809]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
   Bacteriophage phi-42 integrase (int) gene, complete cds
   gi|437115|gb|U01872|U01872 [437115]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 2 protein links, or 3 nucleotide neighbors)
X94423
  Staphylococcus aureus bacteriophage phi-42 DNA with ORFs (restriction modification system)
  gi|1771597|emb|X94423|SARMS [1771597]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 1 nucleotide neighbor)
M27965
  Bacteriophage L54a (from S.aureus) int and xis genes, complete cds
  gi|215096|gb|M27965|L54INTXIS [215096]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, MEDLINE 1 link, 2 protein links, or 3 nucleotide neighbors)
U72397
  Bacteriophage 80 alpha holin and amidase genes, complete cds
  gi|1763241|gb|U72397|B8U72397 [1763241]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)
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AB009866

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Bacteriophage phi PVL proviral DNA, complete sequence
   gi|3341907|dbj|AB009866|AB009866 [3341907]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 63 protein links, or 1 nucleotide neighbor)
 Z47794
   Bacteriophage Cp-1 DNA, complete genome
   gi|2288892|emb|Z47794|BPCP1XX [2288892]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 MEDLINE links, 28 protein links, 1 nucleotide neighbor, or
   I genome link)
 SEG CP7RSIT
   Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat
   gi|166186|gb||SEG CP7RSIT [166186]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
M11635
   Bacteriophage Cp-7 (S.pneumoniae) DNA, 3' inverted terminal repeat
   gi|166185|gb|M11635|CP7RSIT2 [166185]
   (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
M11636
   Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat
   gi|166184|gb|M11636|CP7RSIT1 [166184]
   (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
SEG_CP5RSIT
  Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat
  gi|166181|gb||SEG_CP5RSIT [166181]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
M11633
  Bacteriophage Cp-5 (S.pneumoniae) 3' inverted terminal repeat
  gi|166180|gb|M11633|CP5RSIT2 [166180]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
M11634
  Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat
  gi|166179|gb|M11634|CP5RSIT1 [166179]
  (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
M34780
  Bacteriophage Cp-9 muramidase (cpl9) gene
  gi|166187|gb|M34780|CP9CPL [166187]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
M34652
  Bacteriophage HB-3 amidase (hbl) gene, complete cds
  gi|215055|gb|M34652|HB3HBLA [215055]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
U64984
  Streptococcus pyogenes phage T12 repressor, excisionase (xis), integrase(int) and erythrogenic toxin A precursor (speA) genes.
  complete cds gi[1877426]gb[U40453]SPU40453 [1877426]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 4 protein links, or 22 nucleotide neighbors)
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#### X12375

Phage CP-T1 (Vibrio cholerae) DNA for packaging signal (pac site)
gi|15435|emb|X12375|NCCPPAC [15435]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

# AF087814

Vibrio cholerae filamentous bacteriophage fs-2 DNA, complete genome sequence gi|3702207|dbj|AB002632|AB002632 [3702207] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, or 1 genome link)

#### D83518

Bacteriophage KVP40 gene for major capsid protein precursor, complete cds gi|3046858|dbj|D83518|D83518 [3046858] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

#### AF033322

Bacteriophage PST single-stranded binding protein (gene 32) gene, partial cds, and 5' region gi|2645774|gb|AF033322|AF033322 [2645774] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors)

#### X94331

Bacteriophage L cro, 24, c2, and c1 genes gi|1469213|emb|X94331|BLCRO24C [1469213] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 protein links)

# U82619

Shigella flexneri bacteriophage V glucosyl transferase (gtr), integrase (int) and excisionase (xis) genes, complete cds gi|2465470|gb|U82619|SFU82619 [2465470] (View GenBank report,FASTA report,ASN.1 report,Graphical view,I MEDLINE link, 8 protein links, or 1 nucleotide neighbor)

246 Table 12

NCBI Entrez Nucleotide QUERY

Key words: bacteriophage and lysis

56 citations found (all selected)

# AJ011581

Bacteriophage PS119 lysis genes 13, 19, 15, and packaging gene 3, complete cds gil3676084lemblAJ011581IBPS011581 [3676084] (View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein links, or 1 nucleotide neighbor)

#### AJ011580

Bacteriophage PS34 lysis genes 13, 19, 15, antiterminator gene 23, and packaging gene 3, complete cds gil3676078lemblAJ011580lBPS011580 [3676078] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 2 nucleotide neighbors)

# AJ011579

Bacteriophage PS3 lysis genes 13, 19, 15, and packaging gene 3 gil3676073lemblAJ011579lBPS011579 [3676073] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 protein links, or 1 nucleotide neighbor)

# AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds gil2668751lgblAF034975! [2668751] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

#### U37314

Bacateriophage lambda Rz1 protein precursor (Rz1) gene, complete cds gil1017780[gblU37314|BLU37314 [1017780] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 1 protein link, or 9 nucleotide neighbors)

# U00005

E. coli hflA locus encoding the hflX, hflK and hflC genes, hfq gene, complete cds; miaA gene, partial cds gil436153lgblU00005IECOHFLA [436153]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE

links, 5 protein links, or 8 nucleotide neighbors 3

#### U32222

Bacteriophage 186, complete sequence gil3337249|gblU32222|B1U32222 [3337249] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

# AF064539

Bacteriophage N15, complete genome gil3192683] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

# AF063097

Bacteriophage P2, complete genome gil3139086|gblAF063097|AF063097 [3139086] (View GenBank report,FASTA report,ASN.1 report,Graphical view,21 MEDLINE links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

#### Z97974

Bacteriophage phiadh lys, hol, intG, rad, and tec genes gil2707950lemblZ97974lBPHIADH [2707950]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 9 protein links, or 1 nucleotide neighbor)

# AF059243

Bacteriophage NL95, complete genome gil3088545|gblAF059243|AF059243 [3088545] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 4 protein links, 3 nucleotide neighbors, or 1 genome link)

# AF052431

Bacteriophage M11 A-protein, coat protein, A1-protein, and replicase genes, complete cds gil29812081gblAF0524311 [2981208] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 4 protein links, or 8 nucleotide neighbors)

# Y07739

Staphylococcus phage Twort holTW, plyTW genes gil2764979lemblY07739BPTWGHOLG [2764979] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

# X94331

Bacteriophage L cro, 24, c2, and c1 genes gil1469213lemblX94331lBLCRO24C [1469213] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 protein links)

# X78410

Bacteriophage phiadh holin and lysin genes gil793848lemblX78410LGHOLLYS [793848] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleonde neighbor)

#### X99260

Bacteriophage B103 genomic sequence gil1429229lemblX99260BB103G [1429229] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 17 protein links, or 12 nucleotide neighbors)

# AJ000741

Bacteriophage P1 darA operon gil2462938lemblAJ000741lBPAJ7641 [2462938] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors)

#### X87420

Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators gil1143407]embiX87420BPES18GEN [1143407]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,5 protein links, or 9 nucleotide neighbors)

# L35561

Bacteriophage phi-105 ORFs 1-3 gil532218[gblL355611PH5ORFHTR [532218] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 protein links)

# D10027

Group II RNA coliphage GA genome gil217784ldbjiD10027lPGAXX [217784] (View GenBank report,FASTA report,ASN.1 report,Graphical view,I MEDLINE link, 3 protein links, 5 nucleotide neighbors, or 1 genome link)

# V01128

Bacteriophage phi-X174 (cs70 mutation) complete genome gil15535iemblV01128IPHIX174 [15535]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 11 protein links, or 26 nucleotide neighbors)

# S81763

coat gene...replicase gene [bacteriophage KU1, host=Escherichia coli, group II RNA phage, Genomic RNA, 3 genes, 120 nt] gil1438766[gblS81763|S81763 [1438766] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

# U38906

Bacteriophage r1t integrase, repressor protein (rro), dUTPase, holin and lysin genes, complete cds gil13535171gblU38906IBRU38906 [1353517] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 50 protein links, or 3 nucleotide neighbors)

# X91149

Bacteriophage phi-C31 DNA cos region gil1107473|embiX91149|APHIC31C [1107473] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 6 protein links, or 1 nucleotide neighbor)

#### V00642

phage MS2 genome gil15081lemblV00642lLEMS2X [15081] (View GenBank report,FASTA report,ASN.1 report,Graphical view,8 MEDLINE links, 4 protein links, or 20 nucleotide neighbors)

# V01146

Genome of bacteriophage T7 gil431187lemblV01146!T7CG [431187] (View GenBank report,FASTA report,ASN.1 report,Graphical view,13 MEDLINE links, 60 protein links, 105 nucleotide neighbors, or 1 genome link)

# X78401

Bacteriophage P22 right operon, orf 48, replication genes 18 and 12, nin region genes, ninG phosphatase, late control gene 23, orf 60, complete cds, late control region, start of lysis gene 13 gil512343lemblX78401IPOP22NIN [512343] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 13 protein links, or 4 nucleotide neighbors)

# Y00408

Bacteriophage T4 gene t for lysis protein gil 15368lembi Y00408lMYT4T [15368] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

# **Z26590**

Bacteriophage mv4 lysA and lysB genes gil410500lemblZ26590lMV4LYSAB [410500] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 4 protein links)

#### X07809

Phage phiX174 lysis (E) gene upstream region gil15094lemblX07809lMIPHIXE [15094] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

#### Z34528

Lactococcal bacteriophage c2 lysin gene gil506455lemblZ34528lLBC2LYSIN [506455] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

#### X15031

Bacteriophage fr RNA genome gil15071lembiX15031lLEBFRX [15071] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, 9 nucleotide neighbors, or 1 genome link)

## X80191

Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins gil517237lemblX80191BPP7PR [517237] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 genome link)

#### X85010

Bacteriophage A511 ply511 gene gil853748lemblX85010BPA511PLY [853748] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

#### X85009

Bacteriophage A 500 hol 500 and ply 500 genes gil 853744 lemb 1 X 85009 lBPA 500 PLY [853744] (View Gen Bank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 4 nucleotide neighbors)

#### X85008

Bacteriophage A118 hol118 and ply118 genes gil853740lemblX85008lBPA118PLY [853740] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleoude neighbor)

#### Z35638

Bacteriophage phi-X174 genes for lysis protein and beta-lactamase gil520996lemblZ35638lBPLYSPR [520996] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 516 nucleotide neighbors)

#### J02459

Bacteriophage lambda, complete genome gil215104|gblJ02459|LAMCG [215104] (View GenBank report,FASTA report,ASN.1 report,Graphical view,87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

#### X87674

Bacteriophage P1 lydA & lydB genes gil974763lemblX87674lBACP1LYD [974763] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

#### X87673

Bacteriophage P1 gene 17 gil974761lemblX87673lBACP117 [974761] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

#### M14784

Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis protein and DNA packaging proteins, complete cds gil215810lgblM14784lPT3RE [215810] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, or 10 nucleotide neighbors)

# M11813

Bacteriophage PZA (from B.subtilis), complete genome gil216046lgblM11813IPZACG [216046] (View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 27 protein links, 17 nucleotide neighbors, or 1 genome link)

## M16812

Bacteriophage K3 't' lysis gene, complete cds gil215503lgblM16812lPK3LYST [215503] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

#### J04356

Bacteriophage P22 proteins 15 (complete cds), and 19 (3' end) genes gil215265[gblJ04356IP2215P [215265]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

#### J04343

Bacteriophage JP34 coat and lysis protein genes, complete cds, and replicase protein gene, 5' end gil215076igblJ04343iJP3COLY [215076] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

#### J02482

Bacteriophage phi-X174, complete genome gil216019|gblJ02482IPX1CG [216019]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 23 MEDLINE links, 11 protein links, 26 nucleotide neighbors, or 1 genome link)

#### M99441

Bacteriophage T4 anti-sigma 70 protein (asiA) gene, complete cds and lysis protein, 3' end gil215820igblM99441IPT4ASIA [215820] (View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 2 protein links, or 2 nucleotide neighbors)

#### M65239

Bacteriophage 21 lysis genes S, R, and Rz, complete cds gil215466lgblM65239lPH2LYSGEN [215466] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

# M10637

Phage G4 D/E overlapping gene system, encoding D (morphogenetic) and E (lysis) proteins gil215427|gblM10637|PG4DE [215427] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

# J02454

Bacteriophage G4, complete genome gil215415|gblJ02454|PG4CG [215415] (View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 11 protein links, 20 nucleotide neighbors, or 1 genome link)

#### J02580

Bacteriophage PA-2 (E.coli porcine strain isolate) Rz gene, 5'end; ORF2, outer membrane porin protein (lc) and ORF1 genes, complete cds gil2153661gblJ025801PA2LC [215366] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 4 nucleotide neighbors)

# M14782

Bacillus phage phi-29 head morphogenesis, major head protein, head fiber protein, tail protein, upper collar protein, lower collar protein, pre-neck appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds gil215323lgblM14782lP29LATE2 [215323] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)

#### M10997

Bacteriophage P22 lysis genes 13 and 19, complete cds gil215262lgblM10997lP221319 [215262] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

#### J02467

Bacteriophage MS2, complete genome gil215232|gblJ02467iMS2CG [215232] (View GenBank report,FASTA report,ASN.1 report,Graphical view,8 MEDLINE links, 4 protein links, 20 nucleotide neighbors, or 1 genome link)

#### M14035

Bacteriophage lambda lysis S gene with mutations leading to nonlethality of S in the plasmid pRG1 gil215180[gblM14035lLAMLYS [215180] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors)

# U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein hydrolase (lysB) gene, complete cds gil530796|gblU04309|BPU04309 [530796] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Table 13

# NCBI Entrez Nucleotide QUERY

Key word: holin

51 citations found (all selected)

#### AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds gil2668751gblAF034975 [2668751] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

#### U52961

Staphylococcus aureus holin-like protein LrgA (lrgA) and LrgB (lrgB) genes, complete cds gil1841516]gblU52961|SAU52961 [1841516] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

# U28154

Haemophilus somnus cryptic prophage genes, capsid scaffolding protein gene, partial cds, major capsid protein precursor, endonuclease, capsid completion protein, tail synthesis proteins, holin, and lysozyme genes, complete cds gil1765928|gblU28154|HSU28154 [1765928]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 protein links)

# AF032122

Streptococcus thermophilus bacteriophage Sfi19 central region of genome gil2935682|gblAF032122| [2935682] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

## AF032121

Streptococcus thermophilus bacteriophage Sfi21 central region of genome gil2935667|gblAF032121|AF032121 [2935667]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

#### AF021803

Bacillus subtilis 168 prophage SPbeta N-acetylmuramoyl-L-alanine amidase (blyA), holin-like protein (bhlA), holin-like protein (bhlB), and yolK genes, complete cds; and yolJ gene, partial cds gil2997594lgblAF021803lAF021803 [2997594] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor)

#### AF057033

Streptococcus thermophilus bacteriophage sfill gp502 (orf502), gp284 (orf284), gp129 (orf129), gp193 (orf193), gp119 (orf119), gp348 (orf348), gp53 (orf53), gp113 (orf113), gp104 (orf104), gp114 (orf114), gp128 (orf128), gp168 (orf168), gp117 (orf117), gp105 (orf105), putative minor tail protein (orf1510), putative minor structural protein (orf512), putative minor structural protein (orf1000), gp373 (orf373), gp57 (orf57), putative anti-receptor (orf695), putative minor structural protein (orf669), gp149 (orf149), putative holin (orf141), putative holin (orf87), and lysin (orf288) genes, complete cds gil3320432|gblAF057033|AF057033 [3320432] (View GenBank report,FASTA report,ASN.1 report,Graphical view,25 protein links, or 1 nucleotide neighbor)

#### U32222

Bacteriophage 186, complete sequence gil3337249|gblU32222|B1U32222 [3337249] (View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

# AB009866

Bacteriophage phi PVL proviral DNA, complete sequence gil3341907ldbjlAB009866lAB009866 [3341907] (View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, or 1 nucleotide neighbor)

# AF009630

Bacteriophage bIL170, complete genome gil3282260|gblAF009630|AF009630 [3282260] (View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, 3 nucleotide neighbors, or 1 genome link)

# AF064539

Bacteriophage N15, complete genome

gil3192683|gblAF064539|AF064539 [3192683] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

#### AF063097

Bacteriophage P2, complete genome gil31390861gblAF063097IAF063097 [3139086] (View GenBank report,FASTA report,ASN.1 report,Graphical view,21 MEDLINE links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

## Z97974

Bacteriophage phiadh lys, hol, intG, rad, and tec genes gil2707950lemblZ97974lBPHIADH [2707950] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 9 protein links, or 1 nucleotide neighbor)

#### X95646

Streptococcus thermophilus bacteriophage Sfi21 DNA; lysogeny module, 8141 bp gil2292747lemblX95646lBSFI21LYS [2292747] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 19 protein links, or 3 nucleotide neighbors)

## SEG\_LLHLYSINO

Bacteriophage LL-H structural protein gene, partial cds; minor structural protein gp61 (g57), unknown protein, unknown protein, structural protein (g20), unknown protein, unknown protein, major capsid protein (g34), main tail protein gp19 (g17), holin (hol), muramidase (mur), unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, minor structural protein gp75 (g70), minor structural protein gp89 (g88), minor structural protein gp58 (g71), unknown protein, unknown protein, unknown protein, and unknown protein genes, complete cds gil1004337|gbllSEG\_LLHLYSIN0 [1004337] (View GenBank report,FASTA report,ASN.1 report,Graphical view,4 MEDLINE links, 31 protein links, or 1 nucleotide neighbor)

# M96254

Bacteriophage LL-H holin (hol), muramidase (mur), and unknown protein genes, complete cds gil1004336[gblM96254|LLHLYSIN03 [1004336] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### Y07740

Staphylococcus phage 187 ply187 and hol187 genes gil2764982lemblY07740lBP187PLYH [2764982] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

#### U88974

Streptococcus thermophilus bacteriophage 01205 DNA sequence gil2444080|gblU88974| [2444080] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 57 protein links, or 6 nucleotide neighbors)

#### Z99117

Bacillus subtilis complete genome (section 14 of 21): from 2599451 to 2812870 gil2634966|emb|Z99117|BSUB0014 [2634966] (View GenBank report,FASTA report,ASN.1 report,Graphical view,233 protein links, 51 nucleotide neighbors, or 1 genome link)

# Z99115

Bacillus subtilis complete genome (section 12 of 21): from 2195541 to 2409220 gil2634478|emblZ99115|BSUB0012 [2634478] (View GenBank report,FASTA report,ASN.1 report,Graphical view,244 protein links, 64 nucleotide neighbors, or 1 genome link)

# Z99110

Bacillus subtilis complete genome (section 7 of 21): from 1194391 to 1411140 gil2633472lemblZ99110lBSUB0007 [2633472] (View GenBank report,FASTA report,ASN.1 report,Graphical view,226 protein links, 31 nucleotide neighbors, or 1 genome link)

# X78410

Bacteriophage phiadh holin and lysin genes gil793848lemblX78410lLGHOLLYS [793848] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

# Z93946

Bacteriophage Dp-1 dph and pal genes and 5 open reading frames gil1934760lemblZ93946lBPDP1ORFS [1934760] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 6 protein links)

#### AF011378

Bacteriophage sk1 complete genome gil2392824lgblAF011378lAF011378 [2392824] (View GenBank report,FASTA report,ASN.1 report,Graphical view,54 protein links, 2 nucleotide neighbors, or 1 genome link)

## Z47794

Bacteriophage Cp-1 DNA, complete genome gil2288892lemblZ47794lBPCP1XX [2288892]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 28 protein links, 1 nucleotide neighbor, or 1 genome link)

# L35561

Bacteriophage phi-105 ORFs 1-3 gil532218lgblL35561lPH5ORFHTR [532218] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 protein links)

# D49712

Bacillus licheniformis DNA for ORFs, xpaL2 homologous protein and xpaL1 homologous protein, complete and partial cds gil1514423ldbjlD49712lD49712 [1514423] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 4 protein links)

# X90511

Lactobacillus bacteriophage phigle DNA for Rorf162, Holin, Lysin, and Rorf175 genes gil1926386lemblX90511lLBPHIHOL [1926386] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 protein links, or 1 nucleotide neighbor)

#### X98106

Lactobacillus bacteriophage phigle complete genomic DNA gil1926320lemblX98106lLBPHIG1E [1926320] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

WO 00/32825 PCT/IB99/02040

link, 50 protein links, or 4 nucleotide neighbors)

#### U72397

Bacteriophage 80 alpha holin and amidase genes, complete cds gil1763241|gblU72397|B8U72397 [1763241] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 2 nucleotide neighbors)

# U38906

Bacteriophage r1t integrase, repressor protein (rro), dUTPase, holin and lysin genes, complete cds gil1353517|gblU38906|BRU38906 [1353517] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 50 protein links, or 3 nucleotide neighbors)

#### X91149

Bacteriophage phi-C31 DNA cos region gill 107473|emb|X91149|APHIC31C [1107473] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 6 protein links, or 1 nucleotide neighbor)

# U24159

Bacteriophage HP1 strain HP1c1, complete genome gil1046235gblU24159BHU24159 [1046235] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 41 protein links, 8 nucleotide neighbors, or 1 genome link)

## Z26590

Bacteriophage mv4 lysA and lysB genes gil410500lemblZ26590lMV4LYSAB [410500] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 4 protein links)

# Z70177

B.subtilis DNA (28 kb PBSX/skin element region)
gil1225934|emb|Z70177|BSPBSXSE [1225934]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 32 protein links, or 4 nucleotide neighbors)

Z36941

B.subtilis defective prophage PBSX xhlA, xhlB, and xylA genes gil535793lemblZ36941lBSPBSXXHL [535793]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein links, or 5 nucleotide neighbors)

#### X89234

L.innocua DNA for phagelysin and holin gene gil1134844|emb|X89234|LICPLYHOL [1134844] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

#### X85010

Bacteriophage A511 ply511 gene gil853748lemblX85010lBPA511PLY [853748] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

#### X85009

Bacteriophage A500 hol500 and ply500 genes gil853744lemblX85009lBPA500PLY [853744] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 4 nucleotide neighbors)

# X85008

Bacteriophage A118 hol118 and ply118 genes gil853740lemblX85008lBPA118PLY [853740] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

## L34781

Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and peptidoglycan hydrolase (lytA) gene, partial cds gil511838|gb|L34781|BPHHOLIN [511838] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors)

# U11698

Serratia marcescens SM6 extracellular secretory protein (nucE), putative phage lysozyme (nucD), and transcriptional activator (nucC) genes, complete cds gil509550|gblU11698|SMU11698 [509550] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

link, 3 protein links, or 1 nucleotide neighbor)

# U31763

Serratia marcescens phage-holin analog protein (regA), putative phage lysozyme (regB), and transcriptional activator (regC) genes, complete cds gil965068|gblU31763|SMU31763 [965068] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

#### X87674

Bacteriophage P1 lydA & lydB genes gil974763lemblX87674lBACP1LYD [974763] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

# L48605

Bacteriophage c2 complete genome gil1146276|gblL48605|C2PVCG [1146276] (View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 39 protein links, 3 nucleotide neighbors, or 1 genome link)

# L33769

Bacteriophage bIL67 DNA polymerase subunit (ORF3-5), essential recombination protein (ORF13), lysin (ORF24), minor tail protein (ORF31), terminase subunit (ORF32), holin (ORF37), unknown protein (ORF 1-2,6-12,14-23,25-30,33-36), complete genome gil522252lgblL33769lL67CG [522252] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 37 protein links, 2 nucleotide neighbors, or 1 genome link)

#### L31348

Bacteriophage Tuc2009 integrase (int) gene, complete cds; lysin (lys) gene, 3' end gil508612lgblL31348lTU2INT [508612] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 3 nucleotide neighbors)

# L31364

Bacteriophage Tuc2009 holin (S) gene, complete cds; lysin (lys) gene, complete cds gil496281|gblL31364|TU2SLYS [496281]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

# L31366

Bacteriophage Tuc2009 structural protein (mp2) gene, complete cds gil496278|gblL31366|TU2MP2A [496278] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

# L31365

Bacteriophage Tuc2009 structural protein (mp1) gene, complete cds gil496276|gblL31365|TU2MP1A [496276] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

# U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein hydrolase (lysB) gene, complete cds gil530796|gblU04309|BPU04309 [530796] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

#### Table 14

NCBI Entrez Nucleotide QUERY Key word: bacteriophage and kil 5 citations found (all selected)

#### AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds gil2668751|gblAF034975| [2668751] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

# X15637

Bacteriophage P22 P(L) operon encompassing ral, 17, kil and arf genes gil15646lemblX15637lPOP22PL [15646] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 7 protein links, or 2 nucleotide neighbors)

# J02459

Bacteriophage lambda, complete genome gil215104lgblJ02459lLAMCG [215104] (View GenBank report,FASTA report,ASN.1 report,Graphical view,87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

#### M64097

Bacteriophage Mu left end gil215543|gblM64097|PMULEFTEN [215543] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 39 protein links, or 15 nucleotide neighbors)

# M18902

Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete cds gil166191|gblM18902|D18KIL [166191]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

Table 15

U77328	V01282	U11787	U93688	A47599	D21131	U76864	U38428
AF151117	AF121672	U11786	U93687	A47598	D30690	U76863	U66665
AF151218	AF072726	U11785	AJ224764	A47597	D14711	U76862	U66664
AF146368	AF115379	U11784	AF064774	A47596	D90119	U76861	U66663
AF144661	AF034153	U11783	AF064773	A47595	D00730	U76860	X87104
AF132117	AF029244	U11782	Y14370	A47594	D83357	U76859	X87105
Y15477	U67965	U11781	AF065394	A44534	D83356	U76858	X89233
Y09928	U96610	U11780	AF062376	A44533	D83355	U76857	M28521
Y09594	U96609	U11779	AF062375	A44529	D83354	U76855	U54636
AF134905	U73027	U11778	AF062374	A44528	D83353	U76854	U46541
AB019536	U73026	U11777	AF062373	A44527	D12572	U76853	L14017
AJ237696	U73025	U11776	AB007500	A44526	D86727	U76852	U60589
AF106851	AF068904	U11775	Y09924	A44525	D86240	U76851	X48003
AF106850	U60050	U11774	U63529	A39696	D67075	U76850	M37889
AF106849	D10907	U11773	AF033191	AF001783	D67074	U76849	V01281
M26321	D10906	AF053772	Y15856	AF001782	U97062	U76848	X97985
AF060191	AF053140	AF053771	AB000439	L77194	U96620	U76847	X00127
AF060190	AB013298	AF029731	AF041467	AF003593	U96619	Y09929	X03286
AF060189	Y16431	AF027155	Y14051	AF003592	Z84573	Y09570	X62282
AF060188	AF076684	AF024571	U82085	X73889	AB001896	X95848	X01645
AF060187	AF076683	U87144	AF026122	X74219	Y07645	Y09428	X16471
AF060186	Y13225	AF086644	AF026121	Y10419	U92441	S76611	X52734
AF060185	AF094826	AJ223781	AF026120	M63177	U91741	S76213	X13290
AF060184	AJ223480	AF076030	AB009635	E08773	U29454	S75707	X66088
AF036324	AF093548	AF044951	AB006796	E07163	U29478	S75706	Z30588
AF036323	AJ005352	AF044906	U39769	E07162	U77374	S75705	X16457
AF053568	AF051916	AF044905	D00184	E07161	L42945	S76270	X00342
AJ132841	Y09927	AF044904	X56628	E07160	U38429	S72497	V01287
Y13766	AF051917	AF044903	AF033018	E07159	U81980	S72488	X61307
AF101234	S77058	AF044902	AF034076	E07158	X55185	S74031	Y00356
AJ133520	S65052	AF044901	D82063	E07157	V01278	S67449	X06603
AJ133495	AF009671	AF044900	D76414	E07156	U31979	U75367	Z93205
AJ132803	U81973	AF044899	U57060	E07155	X91786	U75368	X64172
AB016487	U77308	AF044898	D89066	E03836	U36912	U31175	X72700
AB016431	U20869	AF044897	U85095	E03835	U36911	X53096	X60827
AB015981	U89396	AF044075	U85097	E03526	U36910	X53951	X64389
AB015195	U94706	AF044074	U85096	E02873	U64885	X53952	X62288
AF107307	U41072	AF044073	D42078	E01690	U76872	X03408	X55798
AF079518	U52961	AF044072	AF015929	E00876	U76871	U50629	X58434
AJ223806	U21636	AF044071	D10369	E00203	U76870	U38656	X06627
Y18018	U65000	AF044070	A48955	D83951	U76869	U58139	X12831
Y17795	U48826	AF044069	A48501	D17366	U76868	A31894	X07371
AJ005647	U20503	AF044068	A48500	D42144	U76867	L42943	X02529
AJ005646	U11789	AF044067	A48499	D42143	U76866	U51474	Y00688
AJ005645	U11788	AF044066	A47600	D10489	U76865	U50077	X04121
X59477	X54338	A12915	U51133	M63176	M10500	L01053	M63917
X59478	X51661	A12913	U51132	L11998	M10499	M83994	M58515
X63598	X05815	A12906	X02588	L05004	AH000934	J03947	L10909
X52593	X15574	A12905	X61716	L42764	M10498	J03479	M15067

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X76490	Y07536	A12904	X61719	M32103	M10497	M64724	M92376
X81586	X02166	A12903	X61718	U10927	M18264	M14372	M62650
X72014	Z49245	A12902	X67743	AH003057	J01786	M14371	M32312
X72013	X16298	A12901	X67742	M73535	M33833	M14374	M20393
X71437	Z18852	A12900	X67741	M73536	M32470	M15215	M90536
X62992	X68417	A12899	X67740	U20782	M20270	M36694	M21854
X52594	X68425	A12898	X67738	L37598	J03323	M37915	M36771
X14827	X17679	A12897	U02910	L37597	M33479	M12715	L14020
X13404	X63072	A12896	AH003349	L36472	M94061	J04151	M81736
X17301	X02872	A09523	M11118	L25288	M37888	L22566	U11702
X17688	V01277	A04518	M18086	L25893	M76714	L13379	L19300
X03097	X52543	A04517	U19459	K02687	M17123	L13378	L25372
Z16422	A19943	A04512	U35773	L23109	M97169	L13377	L22565
Z33409	A19942	L41499	U26702	L07778	M81346	L13376	M58516
Z33408	A19941	U19770	U21221	M90056	M90693	L13375	U06462
Z33407	A19940	X53818	U36379	J02615	M25257	L13374	L19298
Z33406	A19939	M20129	U06451	M18970	M25256	M17348	M80252
Z33405	A19938	L43098	U35036	K02985	M25255	M17357	L11530
Z33404	A19937	L43082	U20794	M21136	M25254	M17347	
X75439	A19936	X03216	L25426	M10501	M25253	M28364	
X62587	A17958	X70648	M86227	AH000935	M25252	M21319	

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Table 16

# Phage 44AHJD complete genome sequence. 16668 nucleotides.

1	tccatttct	t tactaaacti	t aaaaatgctg	tgcaacaact	taaccaactt	atctaaccta	ttacatattc	
71	atcaaatac	a aaatttatgi	atctattgac	tttattcaa	aattatgatt	tcaacatata	ataaaattaa	
141		t taaatattci						
211		g taaaaccaga						
281		c agaatcaaci						
351	_	c gaagaagaa	-	-			_	
421		g tattagaaca						
491	tgcaacaac	c acaacaagta	a caacaaacac	: aatcagatgt	aacagaatca	aacaaagaag	ataacgacta	
561	ttcagatga	a gaactagttg	y ataagttaga	tttagattag	gaggaattta	aacatgtatg	agggaaacaa	
631	catgcgttc	t atgatgggta	catcatatga	agattcaaga	ttaaataaac	gaacagaatt	aaatgaaaac	
701	atotcaatt	g atacaaataa	aagtgaagat	agttatggtg	tacaaattca	ttcactttca	aaacaatcat	
771		a cgttgaggag						
841		g tcagctaaat						
		g ataattcaaa		_	_	-		
911								
981	_	t aatcgatatt		_				
1051		t gaagaatacg						
1121		a aacgtaatta						
1191	aattcacat	t aaacaacaat	gatacacgtt	tcaatttcca	aacattagca	gacgcaacta	attacgcttt	
1261	aggtgtatad	c aaaaagaaaa	tttctgatat	taatgtatta	gaagaaaaag	aaatgcgtgc	aatgttagtt	
1331	gattactcat	tgaatcaatt	atccgaaaca	aatgtacgta	aagcaacatc	aaaagaagat	ttagcaagca	
1401	-	a agcaatccta	_		_			
1471		ggacaatata						
1541		tttagatac	_				_	
		tgacgactta						
1611	_			_	_	_	-	
1681	_	ttacgtgcgt				_		
1751		, atgtatctaa						
1821		: tattttggat			-			
1891	attccataac	cctgaatttg:	atgaagttac	acactggatt	cattactatt	catttaaagc	cattagtcca	
1961	ttctttaata	aaattttaat	tactgaccaa	gatgtaaatc	caaaaccaga	ggaagaatta	caagaataaa	٠.
2031	aggagcgtaa	aatatgaaca	acgataaaag	aggtttaaac	gttgagttat	caaaggaaat	cagcaaaaga	
2101		atcgcaacag						
2171		: caatcgtgat						
2241		gttggtgaag						
2311		caaattttc						
		acctgactat						
2381								
2451		tttgttgtca						
2521		aattagcaga		-		-		
2591		agaaattaat						
2661		tcacctatgt						
2731	gcattaactg	aaatgaaacg	ggaatatcaa	aacaaaatta	gtgaattaag	taactattta	ggcattaatt	
2801	cattagccgt	tgataaagaa	agcggtgttt	cagacgaaga	ggcaaaaagt	aatcgtggat	ttaccacatc	
2871	aaacaqtaat	atctatttaa	aaggtcgtga	accaattacg	tttttatcaa	agcgttatgg	tttagatatt	
2941	•	acgatgatga		_				
3011		tggctagata						
3081		tgtaaatgat	_	-				
	_	aaagacgttt						
3151								
3221		tactttttaa						
3291		tggcatgcaa						
3361		gttgaaaaat						
3431	actgatgaaa	catcgaatca	aaatgctaca	tctttagaca	attcaactgg	catgactgca	aacagaaacg	
3501		attaccacaa						
3571	taatacgatt	gataacggta	aaactgtgaa	taaatcgagt	aacgaaagta	atcaaaacgc	aaaacgtaat	
3641	caaaatcaaa	aaggtaatgc	aaaaggtaca	caattcacta	agcagtattt	aattgataat	attgataaag	
3711		aagaaagaaa						
3781		ggcatataat						
3851		agagaacgtt						
		atatttcaag						
3921								
3991		aaaacgtttc						
4061		tggttaaacg						
4131		cggcatttgc						_
4201	ttaaagactt	aattaaagat	attgaccgtt	tcgttaatgg	gtttgaatta	aatgagcttg	aaccaaagtt.	-
4271	tgtgatgggc	tttggtggta	ttcgcaacgc	agttaaccaa	tctattaata	ttgataaaga	aacagatcac	
4341	atgtactcta	cacaatccga	ttctcaaaaa	cctgaaggtt	tttggataaa	taaattaaca	cctagtggtg	
4411	acttaatttc	aagcatgcgt	attotacado	gtggtcatgg	tacaacaatc	ggattagaac	qtcaatccaa	
4481		aaaatctggt						
4461 4551	tatatatta	atttagaaga	ggctaeagg	ttaacagatt	atacaccaca	gtcactttta	aacaaacaca	
		gttaattgat						
4621	tatttatate	gcagacgtaa	gaagtaaatg	tastastat	maaaaamar.		taattaaaaa	
4691	ccgcccaaga	gcagacgtaa	aaaa.cacac	racaarara	Juaaaayaad	-yacaattya	caaccayaa	

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aacaatgata atcgttggat gcaaggcatt gctgttgatg gtgatgattt atactggtta agtggtaaca 4761 gttcagttaa ttcacatgtt caaatcggta aatattcatt aacaacaggt caaaagattt atgattatcc 4831 atttaagtta toatatoaag acggtattaa tttcccacgt gataacttta aagagcctga gggtatttgc 4901 atttatacaa atccaaaaac aaaacgtaaa tcgttattac ttgctatgac aaacggcggt ggtggaaaac 4971 gtttccataa tttatatggt ttcttccaac ttggtgagta tgaacacttt gaagcattac gcgcaagagg 5041 ttcacaaaac tataaattaa caaaagacga cggtcgtgca ttatctattc cagaccatat cgacgattta 5111 aatgacttaa cgcaagctgg tttttattat attgacgggg gtactgcaga aaaacttaag aatatgccaa tgaatggtag caagcgtata attgacgctg gttgtttcat taatgtatac cctacaacac aaacattagg 5181 5251 tacggttcaa gaattaacac gtttctcaac aggtcgtaaa atggttaaaa tggtgcgtgg tatgacttta 5321 gacgtattta cgttaaaatg ggattatgga ttatggacaa caatcaaaac tgacgcacca tatcaagaat 5391 atttggaagc aagtcaatac aataactgga ttgcttatgt aacaacagct ggtgagtatt acattacagg 5461 taaccaaatg gaattattta gagacgcgcc agaagaaatt aaaaaagtgg gtgcatggtt acgtgtgtca 5531 agtggtaacg cagtcggtga agtaagacaa acattagagg ctaatatatc ggaatataaa gaattcttca 5601 gtaatgttaa tgcggaaaca aaacatcgtg aatatggttg ggtagcaaaa catcaaaaat aggagtgata 5671 taaatgaaat cacaacaaca agcaaaagaa tggatatata agcatgaggg ggcaggtgtt gactttgatg 5741 gtgcatatgg atttcaatgt atggacttat cagttgctta tgtgtattac attactgacg gtaaagttcg 5811 catgtggggt aatgctaaag acgcgataaa taatgacttt aaaggtttag cgacggtgta taaaaataca 5881 cogagettta aaceteaatt aggggaegtt getgtatata caaatggaca atatggacat atteaatgtg 5951 tgttaagtgg aaatcttgat tattatacat gcttagaaca aaactggtta ggcggcggtt ttgacggttg 6021 ggaaaaagca accattagaa cacattatta tgacggtgta actcacttta ttagacctaa attttcaggt 6091 6161 catattatag aaatgaaaat ggtacattta catgtggttt tttaccaata tttgcacgtg tcggtagtcc 6231 aaaattatca gaacctaatg gctattggtt ccaaccaaac ggttatacac catataacga agtttgttta 6301 tcagatggtt acgtatggat tggttataac tggcaaggca cacgttatta tttaccagtg cgccaatgga 6371 atggaaaaac aggtaatagt tacagtgttg gtattccttg gggggtgttc tcataatggg tattttagcc 6441 tttttttttg aatttagttg 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aaagttagta tcacaatcaa 9031 aacaagctgg acaaccgtct tggtatgacg caggtaacat cgtccacttt gtaccacaag acgtacaaag 9101 aaaaggtaat gcagattttg caaaaaatat gaaagcaggt acaattggac gtgcatatat tccattaaca 9171 gcagctgcta cttgggcggc atattatcct ttaggtttga aagcatcata taacaaagta caaaactatg 9241 gtaatccatt tttagacggt gcgaatacta ttctagcttg gggtggtaaa ttagacggta aaggtggatc 9311 acctagtgat tcgtctgaca gtggtagtag tggtgacagt ggtagttcac tactcgcttt agcaaaacaa 9381 9451 gccatgcaag aattattaaa aaaaatacaa gacgcattac aatgggacgt tcatagtatt ggtagtgata aattttttag taatgattat tttacattag aaaaaacatt taacaacaca tatcatatta aaatgacgat... 9521 tggtttactt gattcattaa aaaaactgat tgatagcgtt caagtagata gtgggagtag tagttctaat 9591 cctactgatg atgacggaga ccataaacca attagtggta aatcagtcaa gccaaatgga aaaagtggtc 9661 gtgtgattgg tggtaactgg acatatgcac agttaccaga aaaatataaa aaagcaattg gtgtaccttt 9731 attcaaaaaa gaatacttat acaaaccagg taacatattt cctcaaacgg gtaatgcagg acaatgtaca 9801 9871 gaattaacat gggcgtatat gtcacaacta catggtaaaa gacaacctac cgacgacggt caaataacaa acggtcagcg tgtatggtac gtctataaaa agttaggtgc aaaaacaaca cataatccaa cagtaggtta 9941 tggtttctct agtaaaccac catacttaca agcaactgca tatggtattg gtcacacagg tgttgttgta 10011

gragtttttg aagatggttc gtttttagtt graaactata atgtaccacc atatgttgca cratcacgtg 10081 10151 tggtattgta tacactcatt aatggcgtac caaataatgc tggtgataat attgtattct ttagtggtat 10221 tqcttaatta actatgctat aatgaacaca tgctagtaat gctagtaaat aaaatacaaa acataatcaa ttttcgtaca catttttcat gttatctcaa aaagaaaagg agactgttat tttaacagtt gcctttttt 10291 atttcatcat gttcacgttt taatatatgc aaatcagatt tgttatgtac tgaacgttca actggaaata 10361 agtogttaag tgaaaatgaa cogatgtcac tttcaatata aagaatatca tcaaattgac tatggtcgaa 10431 attttcccta gcgtctttta atataaattc acgtttcata ttaagttcat cagtaaaata ttcatcatat 10501 acattaccac atacaatttc agttttagac ggatatatcg atattgtacc ttgctcatta tagatacttt 10571 tattqttttc aataatggca ccgtcaaaga attgttcacg tacaaaggtt tcaaaatcga cgcttgtatc 10641 aaaggegttt tteggtatae cageagaage aattttaate tttecattea etteatatge atatttetta 10711 tgattcagta caaacatctt atctatctgt tcgttttcaa tatcccattt acctaaggct atcgggtcga 10781 ataaactggg gttcaataag ggtttaacaa cggatttcat atacaaacta tcagtatcgc aataaataaa 10851 attgtegtea attteaettt eegttaagta ttggaaagga accaataagt tatacaatga aegtgatgtg 10921 acasatqtaq aqaataatat attacgttca gtgtttttgt aaccgttaat gatattgtat agttcattgt 10991 tatcatctaa acggaataag ttaaaatgtg aacgtaatgc aggtatgcca tataatccat ttaaaacgac 11061 tttagataac ataacctcct catttgagta tgggtgttcg ttgatatcat cagtaatgtg atagtcgtaa 11131 ggtgatgtca tattgatttt gttttttaac ttaccttgtg ttttaataaa atagttttga aaaataatat 11201 cacgtgcatg aaagtattca cattcatata taacaaacga attaacacgt atatgcatgc aatcaatacc 11271 cgtaatgtct tgaatcattc ttaatgtatt tgtattgata ttaacgtaat cattatcatt attatagtat 11341 tttacaatca tttgacgtaa tacacgtgat ttaattttaa ttaataaatc atcgttaaat acatctttat 11411 caatcttata taatgaaaaa taattgtcat catctaaaaa agtagggatt aacgttggtt ctgaatagtg 11481 ttcgtaaaag tataaccatg ttggaatttt ttcatgatac atcacataag gataactcga attgatgtca 11551 atagaaaaac aaggeteate aattagtttg tttatgtatt tggtgttata catatttaaa ccaccacgat 11621 agaatgattt aatatagtca taaaaattca tatcatggaa atgataatgt gtataagata ttttaatatc 11691 ttgatattgg ttgagtaact gaaaacgtgt catttcatta ttcaagtaag attccataat attcaatgaa 11761 aatgttaatt tgttatagte aaaatttgga aatatatcac tataatgaat atggcacata cctaatataa 11831 tcacgtcatt atgaatgtat gtaagttgtt caggtgtgag ttttgcaaaa catttcacag catagtcata 11901 ggcttcacta tcattcatat cattatcttt atcaaaaatc gtataattaa aatctgtttt aagttgtgat 11971 totgttaaat aaccaccatc aagtaattto ttacctaatg ttgcaattga tgtattggtt ttcataaagt 12041 tatcaataat attaaattta aaaccattta aaaacattgt taaatctaaa ttgattgaag atttaacacg 12111 tttttctaaa attacatttt gatttttggc taaaatagta gcctctttca tttttaatgt gtgttcattt 12181 tettetqeag attttaaata tatatttteg egtgtaatat tatcaaaata acgeatggtg tetttaagta 12251 aaaaatgatt atcgtattta ttacagttat gtgcaatcat gataatatct gtttttgatt ttgtgattgt 12321 atcacgtett ttcacatacg tataaaatge gtcataaaaa gattegaaac teggaaatac ttcaacatca 12391 atttcataac cattaaacca accaattgct acagaataag taacgttttt atatttggtt ggtttttttc 12461 gtccgttaac tttattgtac gctaatgttt ctatatccca gtataaaatc attcgacgtt catgtttatg 12531 atattgcatg cattctagta atcccataat cttacacacc ttttataagc catattgttt cattagatac 12601 tttttcgtat tctctatata gttatcttcg tatatttttt cttttctttc aaactcactc atatttttct 12671 tcatttcatt ttttatatga aattttataa ttttattcat atctaaatat aaatatctat cattatcaac 12741 cacqtaattt ttaqaqtaaq cattqtcaaa atgtaaattg cttggattgt agtaataacg ttccatgttt 12811 totttataaa acatatcato acgtaaatag gtaacatgat tgtctatato cotaatttta gtacaaaatt 12881 catattgttt tgtatatggt acaacgataa tatttgtcat aaaagtagtt acattataca tgactttaat 12951 atatttatca tcagttttga tatagaagaa atcaccgttt tgattgatgt gatttcttaa attatcatcc 13021 gccaaattat attcgttaaa ttcaaattct ccagttgtca tagcgtcgtc atttgaatta aacgcacgtg 13091 tgttacgttt ttcattcacg taatcgtttc gtcgcatttc taaaaaaatg tttttgtaaa gtcttgatgt 13161 attcatttta tgcttttgta ataaattgta tatatttaaa ttggataata taggacttga aaagttgact 13231 gcattaccta gtaaaaacat tttagggaat ccaatataat caacgttacc atggttacgg tcgattgatt 13301 catatattgt ttttaactta toccactcat caattaaata atcatottca agtgctaaaa actcatcata 13371 tataataata ggatagtgtt ttaaaaagtt agaatgatat tttaaatcag tggcactatt caaatctgta 13441 atcacaccaa tttctttatc ttgatagata atagctaaat agtccctagc acttctgaac gtgacacgtt 13511 13581 ttgatttaaa tagtggattt tcatctatga tttcttcaat aaaatcacgg taagcgtcac gtaatgtata atgacgtgat aataaagtaa attttatatc aagtttaata gctaaataaa taaaaaatga aacatagttg 13651 aacgattttc catcagaacg gtttgaaata gatatataat aatctatatc atcattcata agttcatcaa 13721 ctaattctat ttgattatac ttatctggga ttttttttct gacatgattg acagcatttt gataatctct 13791 taccatqtct aaacqatttt gttttaccat gtttttgctc cttgtaatag tttatgatgt cgtttacagt 13861 gttaaattta ttcgtcaaat gttgcataat ataaaaagtt atacctcaca tcttcatcat caatatttgt 13931 cactggtcta totgatttac caatttottt atataaagta togatttott taatatattt atacattgaa 14001 gaattattat ttttagettg taaattatat aaagegtatt tatgettttt agegtttta ttattagaat 14071 14141 agcatttaca tatgatacgt ttctttcttt aggaaaatag ggcagatgtg caaaatgttt ccatgtgtca 14211 14281 ctttttcttg ctcttttcta gcttctcttt ctttttcca tctatccatt tcagacgtat gtctaaccaa 14351 tgttatcaac ctccatataa agcataaata accattaaaa agataatata gaatataatc aatgtagtga 14421 14491 ataaaacacc aaatgacacg cgtatatgca gtgtcataag tatgataagt gtaattaaaa atgctaaaag 14561 gaaaacaatg gctatgttta ataggttatt catggtcaat cactttccca ttatcgtata tgactttgtt 14631 ttgataaata atcattaatt cgctttcaag aggtttatca aaatttgata atacgtcgtc aattgtaacg 14701 tttaataaaa tttctcttat taattcatta cttaaataat ttctataata aaatacaagt atattaaaaa catgittitt aatatcaatg togatatota acgtaaataa cictittica atticaaaat catcatatig 14771 tttgtcaaac tcaatataca catcacccat atttatttt actatacatt ttttattaga tgaagtaaat ... 14841 ttttcaaatt tatcattata ataatctcta tttgttaaaa ggtaataaat taaattattt aatctaaazg 14911 14981 tagttttaat tttcattttt atatctcctt aatgtattct atgatatacg cgtatttttt agtgaacagg ttatattcat aatatgaata tacaacttta gogtcatata aatcttcaaa cattgagatt tgatgtggaa 15051 aatgteettt aateteateg caatataata atacegtttt gtatttaegt teeatttaaa caceteataa 15121 aaaataqqqq ataaqtatcc cctatgaaat tgtattaaaa tgatacttga ccaaaattga ttgagtaacc 15191 tttttgacct tttttgtttt catattcata aattgtgaat tgaacttctc cagcattgat aatgtcaaca 15261 acgtecteat etgeteteat ttetttaatt aattetgtta agtggttegg taagtttaeg ttatagteat 15331

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15401	cagtgacgat	aacaccttgt	tcaccgaatt	ttgattcttt	gtttgtgaat	aatgctctaa	cgatatactc
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15541	aatctcgcta	atgtgttttg	gtgtcttgat	aaaatatctt	ttacgtttgt	cattttattt	ctcctcttat
15611	ttaaattatt	tgctttctgc	aattgcgatt	tgtagtaaat	cattgtaata	aacttgaatt	gttttcgttg
15681	tacatataat	ggacaatagt	ttacatgtgt	ctggtaataa	ttcttttgct	tgtgttttgg	ttaaatgata
15751	ctcqtqaaqt	ggtaaaaatt	cctcaatgta	ttcattatca	tcatctaagt	aatgaagtat	ataacctttg
15821	acacqtaaqq	taacaatqtc	gtcaactttc	attattatat	cactcctttc	taaaaaacgt	aaacgttata
15891	cqtttcataa	aatcctttat	gcatattcca	ttgttctatt	gggtcatcac	cagcaatata	agacaatatt
15961	gattetggtt	tagtttcgtt	gtttagttca	tcatttaaga	attgaacaac	agaactatta	tagtttaata
16031	atagttgttg	gcaagccgat	aataagttaa	ttgcattgtc	aaatgtataa	gctggattcc	attgaatcag
16101	tttattqaat	agttgcaaca	tttcagtata	ggcttgtcct	ttttcttctg	gtgcattatc	aacattaacc
16171	attattatca	cttcctaata	aagttgaaat	tacgcgtaaa	acagaattat	gatttaaatc	ttcaatttca
16241	tcaatgtcaa	catcataaaa	tgaaatttca	ttttctgttc	tatcaaataa	cgctatacat	aaacttccat
16311	tcttaaaacq	aaaaacatgc	ttcaactcaa	tgttttttgt	ttcattttcc	atttttgtta	ctccttgttt
16381	tgattacata	cttagtatag	caaacgttta	aaagttttgt	caatagtttt	tcttaaaaaa	gtttaaataa
16451	ttttaaaact	actatttaat	agaagaaata	agattttaag	ttcaaatcat	aattttgaat	aaaagtcaat
16521	agatacataa	attttqtatt	tgatgaatat	gtaataggtt	agataagttg	gttaagttgt	tgcacagtat
16591	ttttaagttt	agtaaagaaa	tgataagtaa	atttataagt	tttgatttgt	ataatcgttt	attttaaacc
16661	ggtggggt			_	_		

Table 17

# Phage 44AHJD ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	44AHJDORF001	-1	1034212627	761	DNA polymerase;
2	44AHJDORF002	3	37895732	647	Techoic acid; Staph;
3	44AHJDORF003	2	66268389	587	Tail;
4	44AHJDORF004	1	876410227	487	Serine protease motif;
5	44AHJDORF005	-1	1264313890	415	
6	44AHJDORF006	2	8032029	408	
7	44AHJDORF007	1	20443027	327	Upper collar;
8	44AHJDORF008	2	3020,.3775	251	Lower collar;
9	44AHJDORF009	2	57446496	250	Amidase: Staph;
10	44AHJDORF010	-2	1393814420	160	
11	44AHJDORF012	3	83918813	140	Holin;
12	44AHJDORF013	-2	1458614996	136	7.10.11.1
13	44AHJDORF113	1	199600	133	
14	44AHJDORF011	-2	1522515593	122	- <del> </del>
15	44AHJDORF114	-2	1587016172	100	
16	44AHJDORF014	3	62436521	92	<del></del>
17	44AHJDORF015	1 1	1540315645	80	
18	44AHJDORF016	-1	1561615852	78	
19				73	
20	44AHJDORF017 44AHJDORF018	-2	1053610757 8861098	70	<u> </u>
				68	
21 22	44AHJDORF019	-2 -1	96309836	65	
	44AHJDORF121		1616516362	62	
23	44AHJDORF020	2	1386514053	60	<u> </u>
24 25	44AHJDORF123	-2	614796	60	
	44AHJDORF021		56345816	59	
26	44AHJDORF023	-2	63156494	58	
27	44AHJDORF024	1	1427514451	<u> </u>	
28	44AHJDORF025	-3	1499915175	58	
29	44AHJDORF026	-3	1442614593	55	
30	44AHJDORF027	1	1291613080	54	
31	44AHJDORF029	-1	1501915183	54	ļ
32	44AHJDORF028	-3	90719235	54	
33	44AHJDORF030	3	1448714648	53	
34	44AHJDORF031	2	1103911191	50	<u> </u>
35	44AHJDORF135	3	693842	49	
36	44AHJDORF033	-1	36463795	49	
37	44AHJDORF032	-2	93069455	49	
38	44AHJDORF034	-3	1400014146	48	
39	44AHJDORF035	-3	1381113957	48	
40	44AHJDORF036	-3	1001910165	48	
41	44AHJDORF022	-3	84688611	47	
42	44AHJDORF037	1	1478814931	47	
43	44AHJDORF038	-2	35283671	47	
14	44AHJDORF039	3	17431883	46	
15	44AHJDORF040	2	97409877	45	
16	44AHJDORF041	2	1583615973	45	
17	44AHJDORF042	-1	50145151	45	
18	44AHJDORF043	-1	4402.,4539	45	
19	44AHJDORF044	-2	1278312917	44	
0	44AHJDORF149	-2	639770	43	
i1	44AHJDORF046	1	48915019	42	
2	44AHJDORF047	1	1191112039	42	
3	44AHJDORF045	2	1065510783	42	<u> </u>
4	44AHJDORF048	-3	1521215340	42	
5	44AHJDORF049	3	57845909	41	
6	44AHJDORF050	3	1315813283	41	
7	44AHJDORF051	-2	1094411066	40	
8	44AHJDORF052	-3	1421614338	40	
9	44AHJDORF053	3	33483467	39	
ō	44AHJDORF054	3	75517670	39	
1	44AHJDORF055	3	1570515821	38	
2	44AHJDORF056	1	55125625	37	
3	44AHJDORF057	2	1012110231	36	
- 1	7 17 11 10 D Q 1 11 O O 1	3	1076710877	36	

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65	44AHJDORF164	-1	592702	36	
66	44AHJDORF059	-2	82508360	36	
67	44AHJDORF060	-2	61476257	36	
68	44AHJDORF061	2	1555115658	35	
69	44AHJDORF062	1	42854389	34	
70	44AHJDORF063	-3	93839487	34	
71	44AHJDORF065	1	50295130	33	
72	44AHJDORF064	2	26092710	33	
73	44AHJDORF066	-2	1038010481	33	

\_\_\_\_

#### Table 18

# Predicted amino acid sequences

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K V N G R K P T K Y K N V T Y S V A I G W F N G Y E I
12459 gatgttgaagtatttccgagtttcgaatctttttatgacgcattttatacgtatgtgaaaagacgtgatacaatcacaaaatca
       D V E V F P S F E S F Y D A F Y T Y V K R R D T I T K S
      12375
       K T D I I M I A H N C N K Y D N H F L L K D T M R Y F D
12291 aatattacacgcgaaaatatatatttaaaatctgcagaagaaaatgaacacacattaaaaatgaaagaggctactattttagcc
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113
       12207
       KN Q N V I L E K R V K S S I N L D L T M F L N G F K F
12123 aatattattgataactttatgaaaaccaatacatcaattgcaacattaggtaagaaattacttgatggtggttatttaacagaa
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169
      12039
       SQLKTDFNYTIFDKDNDMNDSEAYDYAV
197
K C F A K L T P E Q L T Y I H N D V I I L G M C H I H Y
225
       agtgatatatttccaaattttgactataacaattaacattttcattgaatattatggaatcttacttgaataatgaaatgaca
S D I F P N F D Y N K L T F S L N I M E S Y L N N E M T
11871
       {\tt cgttttcagttactcaaccaatatcaagatattaaaatatcttatacacattatcatttccatgatatgaatttttatgactat}
11787
       R F Q L L N Q Y Q D I K I S Y T H Y H F H D M N F Y D Y
281
       11703
        I K S F Y R G G L N M Y N T K Y I N K L I D E P C F S I
309
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11619
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337
       acgttaateeectaettttttagatgatgaeaattattttteattatataagattgataaagatgtatttaaegatgatttatta
11535
        T L I P T F L D D D N Y F S L Y K I D K D V F N D D L L
365
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11451
393
       a catta agaa t gatt caaga catta c g g g tatt g att g cat g cat at a c g t g t tatt c g t t t g t t a t a t g a t g cat a c g t g t a t a t g cat a c g c g cat a t g cat a c g c g cat a t g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a c g cat a
11367
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421
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10947
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561
       10863
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10779
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10695
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645
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10611
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673
       {\tt tttatattaaaagacgctagagaaaatttcgaccatagtcaatttgatgatattctttatattgaaagtgacatcggttcattt}
10527
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10443
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729
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10359
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757
44AHJDORF002
        atggcatataatgaaaacgattttaaatattttgatgacattcgtccatttttagacgaaatttataaaacgagagaacgttat
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        3873
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29
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3957
57
        ccagagcaagcgaaagacttatttagaggttggttaaacgacggtacgattgacagtattattcatgacgagtttaaaaaatat
4041
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        4125
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113
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4209
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197

225 7382

253

7466

281

7550

309

337

365 7802

393

7886

421

449

8054

477 8138

7970

7718

7634

WO 00/32825 PCT/IB99/02040

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533
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8389
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561
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29
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9016
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9100
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9268
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169
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9352
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197
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225
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9520
253
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9604
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281
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    PISGKSVKPNGKSGRVIGGNWTYAQLPE
9688
309
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9772
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337
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421
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10192
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477
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57
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13638
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13554
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13470
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141
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13386
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13302
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197
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12714
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44AHJDORF006
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1055
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197
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225
1559
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253
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1727
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337
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1895
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1979
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2128
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2968
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3272
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3356
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113
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3440
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3524
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3692
3775
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225
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44AHJDORF009

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5996
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6080
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6164
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6248
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6332
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6416
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225
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14420
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14336
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14168
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8475
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57
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8643
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1
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283
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367
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         INWGSIRV *
44AHJDORF164
         at {\tt gttttcatttaatcttgttcgtttatttaatcttgaatcttcatatgatgtacccatcatagaacgcat {\tt gttttccctcatatgatgtacccatcatagaacgcat {\tt gtttccctcatatgatgtacccatcatagaacgcat {\tt gtttccctcatatgatgtacccatcatagaacgcat {\tt gtttccctcatatgatgtacccatcatagaacgcat {\tt gtttccctcatatgatgtacccatcatagaacgcat {\tt gtttccctcatatagaacgcat {\tt gtttccctcatatagaacgcat {\tt gttccctcatatagaacgcat {\tt gtttccctcatatagaacgcat {\tt gttccctcatatagaacgcat {\tt gttccctcatatagaacgcatcatcatagaacgcat {\tt gttccctcatatagaacgcatcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcatcatagaacgcat
702
         MFSFNSVRLFNLESSYDVPIIERMLFPS
1
         tacatgtttaaattcctcctaatctaa 592
618
29
         YMFKFLLI *
44AHJDORF059
         arggattttgtaacattggattacctgaaccgtcattatgccaaaatcttacaccagattctaaaattgcttttaattgctcca
         M D F V T L D Y L N R H Y A K I L H Q I L K L L H I V P
         ttaacatggggtcgatgtcacgtatag 8250
8276
         L T W G R C H V *
29
44AHJDORF060
         at \texttt{gtaccattttcatttctata} at \texttt{at} \texttt{gtgccgtattggtttcgtttccattttcca} \\ \texttt{at} \texttt{gtatttacttttgatgtttctaatg}
6257
         MYHFHFYNMCRIGFVSIFQMYLLLMFLM
```

```
6173
       ctttgctattactacctgaaaatttag 6147
       LCYYYLKI *
44AHJDORF061
{\tt 15551} \quad {\tt atgtgttttggtgtcttgataaaatatcttttacgtttgtcattttatttctcctcttatttaaattatttgctttctgcaatt}
       MCFGVLIKYLLRLSFYFSSYLNYLLSAI
1
     gcgatttgtagtaaatcattgtaa 15658
15635
29
       AICSKSL *
44AHJDORF062
      gtggtattcgcaacgcagttaaccaatctattaatattgataaagaaacaaatcacatgtactctacacaatccgattctcaaa
4285
       V V F A T Q L T N L L I L I K K Q I T C T L H N P I L K
1
       aacctgaaggtttttggataa 4389
4369
29
       N L K V F G *
44AHJDORF063
9487
      atgcgtcttgtatttttttttaataattcttgcatggcttgttttgctaaagcgagtagtgaactaccactgtcaccactactac
       M R L V F F L I I L A W L V L L K R V V N Y H C H H Y Y cactgtcagacgaatcactag 9383
1
9403
29
       H C Q T N H *
44AHJDORF065
      gtggtggaaaacgtttccataatttatatggtttcttccaacttggtgagtatgaacactttgaagcattacgcgcaagaggtt
VVBNVSIIYMVSSNLVSMNTLKHYAQEV
1
       cacaaaactataaattaa 5130
5113
      HKTIN *
29
44AHJDORF064
      atgacgagtcaatcaatcaacttgtgtccgaaatatataacggtgcaccatttgttaaaatgtcacctatgtttaatgcagatg
2609
       M T S Q S I N L C P K Y I T V H H L L K C H L C L M Q M
      acgatatcattgatttaa 2710
2693
29
       TISLI *
44AHJDORF066
atgatattetttatattgaaagtgacateggtteatttteaettaaegaettattteeagttgaaegtteagtaeataaeaaat

M I F F I L K V T S V H F H L T T Y F Q L N V Q Y I T N
10397 ctgatttgcatatattaa 10380
      LICIY *
29
```

# Table 19

# Sequence similarities between ORFs 44AHJD and public databases

```
Phage: 44AHJD
Database: nr
Query= sid|110871|lan|44AHJDORF001 Phage 44AHJD ORF|10342-12627|-1
           (761 letters)
gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0...
                                                                                   55 1e-06
gi|1072656|pir||S51275 DNA polymerase - phage CP-1 >gi|836593|e...
                                                                                        6e-06
                                                                                   53
gi|1429230|emb|CAA67649| (X99260) DNA polymerase [Bacteriophage...
gi|1572479|emb|CAA65712| (X96987) DNA polymerase [Bacteriophage...
gi|118851|sp|P06950|DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...
                                                                                   49 le-04
                                                                                   46
                                                                                        0.001
                                                                                        0.002
gi|2435429 (AF012250) unassigned reading frame (possible DNA po...
                                                                                   45
                                                                                        0.002
gi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum po...
                                                                                   45 0.002
gi|4877819|gb|AAD31446.1| (AP133505) DNA polymerase [Neurospora...
                                                                                        0.004
                                                                                   44
gi|461962|sp|P33537|DPOM_NEUCR PROBABLE DNA POLYMERASE >gi|2833...
                                                                                   44 0.004
gi|2499511|sp|Q12471|6P22_YEAST 6-PHOSPHOFRUCTO-2-KINASE 2 (PHO...
gi|2258375|gb|AAD11909.1| (AF007261) transcription initiation f...
                                                                                   41 0.041
                                                                                   40
                                                                                        0.070
gi|15734|emb|CAA37450| (X53370) DNA polymerase (AA 1-575) [Bact...
                                                                                   39 0.092
Query= sid|110872|lan|44AHJDORF002 Phage 44AHJD ORF|3789-5732|3
           (647 letters)
gi|135273|sp|P27622|TAGC_BACSU TEICHOIC ACID BIOSYNTHESIS PROTE...
                                                                                  112 7e-24
gi|142847 (M64050) DNase inhibitor (Bacillus subtilis)
                                                                                   52 1e-05
                                                                                   39 0.10
gi|4038407 (AF103943) factor C protein precursor [Streptomyces ...
Query= sid|110873|lan|44AHJDORF003 Phage 44AHJD ORF|6626-8389|2
           (587 letters)
gi|138123|sp|P04331|VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                                   92 8e-1B
gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                                        1e-14
gi|1429238|emb|CAA67657| (X99260) tail protein [Bacteriophage B...
gi|215339 (M12456) p9 tail protein [Bacteriophage phi-29] >gi|2...
gi|1181968|emb|CAA87738.1| (Z47794) tail protein [Bacteriophage...
gi|1181970|emb|CAA87740.1| (Z47794) tail protein [Bacteriophage...
                                                                                   78 2e-13
                                                                                   71 2e-11
                                                                                   54 3e-06
                                                                                   42 0.010
Query= sid|110875|lan|44AHJDORF005 Phage 44AHJD ORF|12643-13890|-1
          (415 letters)
gi|3845203 (AE001399) GAP domain protein (cyclic nt signal tran...
                                                                                   52 6e-06
gi|3758843|emb|CAB11128.1| (298551) predicted using hexExon; MA...
                                                                                        5e-05
                                                                                   49
gi|3845297 (AE001421) hypothetical protein [Plasmodium falciparum]
                                                                                   48 le-04
gi|4493936|emb|CAB38972.1| (AL034556) predicted using hexExon; ...
                                                                                   47
                                                                                        2e-04
gi|3845165 (AE001390) hypothetical protein [Plasmodium falciparum]
                                                                                   46 6e-04
Query= sid|110877|lan|44AHJDORF007 Phage 44AHJD ORF|2044-3027|1
           (327 letters)
gi|1181960|emb|CAA87731.1| (Z47794) connector protein [Bacterio...
                                                                                   46 5e-04
gi|1429239|emb|CAA67658| (X99260) upper collar protein [Bacteri...
gi|137915|sp|P07535|VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...
                                                                                   45
                                                                                        8e-04
                                                                                        0.002
                                                                                   44
gi|137914|sp|P04332|VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...
                                                                                   41 0.009
Query= sid|110878|1an|44AHJDORF008 Phage 44AHJD ORF|3020-3775|2
           (251 letters)
gi|4982468|gb|AAD30963.2| (AF118151) SNF1/AMP-activated kinase ...
                                                                                   52 3e-06
gi|1730077|sp|P18160|KYK1_DICDI NON-RECEPTOR TYROSINE KINASE SP...
gi|3758855|emb|CAB11140.1| (298551) predicted using hexExon; MA...
                                                                                        2e-04
                                                                                   46
                                                                                   46 2e-04
gi|585795|sp|P21538|REB1_YEAST DNA-BINDING PROTEIN REB1 (QBP) >...
                                                                                   46
                                                                                        3e-04
gi|172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]
                                                                                   46 3e-04
                                                                                        6e-04 _
gi|2952545 (AF051898) coronin binding protein [Dictyostelium di...
                                                                                   45
gi|535260|emb|CAA82996| (Z30339) STARP antigen [Plasmodium reic...
                                                                                   45 7e-04
                                                                                   44 0.001
gi|1429240|emb|CAA67659| (X99260) lower collar protein [Bacteri...
```

# Query= sid|110879|lan|44AHJDORF009 Phage 44AHJD ORF|5744-6496|2 (250 letters)

(250 lecters)		
gi 2764981 emb CAA69021.1  (Y07739) N-acetylmuramoyl-L-alanine	180	1e-44
qi   113675   sp   P24556   ALYS STAAU AUTOLYSIN (N-ACETYLMURAMOYL-L-AL	118	6e-26
gi 1763243 (U72397) amidase (bacteriophage 80 alpha)	118	6e-26
gi 4574237 gb AAD23962.1 AF106851_1 (AF106851) LytN [Staphyloco	84	9e-16
gi 3767593 dbj BAA33856.1 (AB015195) Lyth (Staphylococcus aureus)	84	9e-16
qi 2764983 emb CAA69022.1  (Y07740) cell wall hydrolase Ply187	77	2e-13
qi 3287732 sp 005156 ALE1 STACP GLYCYL-GLYCINE ENDOPEPTIDASE AL	73	2e-12
gi 79926 pir A25881 lysostaphin precursor - Staphylococcus sim	69	3e-11
qi 126496 sp   P10548 LSTP STAST LYSOSTAPHIN PRECURSOR (GLYCYL-GL	69	3e-11
gi 3287967 sp   P10547   LSTP_STASI LYSOSTAPHIN PRECURSOR (GLYCYL-G	69	3e-11
gi 3341932 dbj BAA31898.1  (AB009866) amidase (peptidoglycan hy	68	6e-11
Query= sid 110882 lan 44AHJDORF012 Phage 44AHJD ORF 8391-8813 3 (140 letters)		
gi 140528 sp P24811 YQXH BACSU HYPOTHETICAL 15.7 KD PROTEIN IN	80	6e-15
gi 4126631 dbj BAA36651.1  (AB016282) ORF45 [bacteriophage phi	76	1e-13
gi 141088 sp P26835 YNGD_CLOPE HYPOTHETICAL 14.9 KD PROTEIN IN	61	4e-09
gi 2293160 (AF008220) YtkC [Bacillus subtilis] >gi 2635548 emb	36	0.099
gi 1181973 emb CAA87743.1  (Z47794) holin protein [Bacteriophag	31	3.3

#### Table 20

# Homolgies between phage 44 AHJD ORFs and proteins in public databases

```
Query= pt|110871 44AHJDORF001 Phage 44AHJD ORF |10342-12627|-1 1
         (761 letters)
>gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0161
          DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2
          >gi|215509 (M33144) DNA polymerase [Bacteriophage M2]
          Length = 572
 Score = 55.4 bits (131), Expect = 1e-06
 Identities = 96/426 (22%), Positives = 159/426 (36%), Gaps = 88/426 (20%)
Query: 229 KLTPEQLTYIHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTR-----FQ 283 ++TPE+ YI ND+ I+ DI +++T + ++ + T F
Sbjct: 154 EITPEEYEYIKNDIEIIARA----LDIQFKQGLDRMTAGSDSLKGFKDILSTKKFNKVFP 209
Query: 284 LLNQYQDIKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYP 343
           L+ D +I
                          + YRGG NKY KIE
Sbjct: 210 KLSLPMDKEI------RKAYRGGFTWLNDKYKEKEIGEGMV-FDVNSLYP 252
Query: 344 YVMYHEKIPTWLYFYEHYSEPTLIPTFLDDDNYFSLYKIDKDVFNDDLLIKIKSRVLRQM 403
            MY +P Y P + + D + LY I + F +L K + +
Sbjct: 253 SQMYSRPLP-----YGAPIVFQGKYEKDEQYPLY-IQRIRFEFEL----KEGYIPTI 299
Query: 404 XXXXXXXXXXXXXXXXXXXXXIRMIQ-DITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
                           + ++ +T +D I+ + +Y EY
Sbjct: 300 QIKKNPFFKGNEYLKNSGVEPVELYLTNVDLELIQEH-YELYNVEYIDGFK-----FRE 352
Query: 463 TQGKLKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPAL 511
            G K+ I+ + H + L+K++LN LYG
Sbjct: 353 KTGLFKDFIDKWTYVKTH------EEGAKKQLAKLMLNSLYGKFASNPDVTGKVPYL 403
Query: 512 RSHFNL-FRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNF 570
                         YK+ + F+T+ + + Q
             +L FR+ D
Sbjct: 404 KDDGSLGFRVGDEE------YKDPVYTPM-GVFITAWARFTTITAAQACY-----DRI 449
Query: 571 IYCDTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKK-----YAYEVNG 625
          IYCDTDS+++ P + + DP LG W E+ + L K
Sbjct: 450 IYCDTDSIHLTGTEVPEIIKDIVDPKKLGYWAHES-TFKRAKYLRQKTYIQDIYVKEVDG 508
Query: 626 KIKIAS 631
          K+K S
Sbjct: 509 KLKECS 514
>gi|1072656|pir||S51275 DNA polymerase - phage CP-1
          >gi|836593|emb|CAA87725.1| (Z47794) DNA polymerase
          [Bacteriophage CP-1]
          Length = 568
 Score = 53.5 bits (126), Expect = 6e-06
Identities = 104/464 (22%), Positives = 169/464 (36%), Gaps = 66/464 (14%)
Query: 230 LTPEQLTYIHNDVIIL--GMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTRFQLLNQ 287
          + PE + YIH DV IL G+ ++Y + F Y + +L +
Sbjct: 152 IKPEWIDYIHVDVAILARGIFAMYYEENFTK--YTSASEALTEFKRIFRKSKRKFRDFFP 209
Query: 288 YQDIKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMY 347
                          D+ + G + K+ + +++ DINS YP M
Sbjct: 210 ILDEKVD------DFCRKHIVGAGRLPTLKHRGRTLNQLIDIYDINSMYPATML 257
Query: 348 HEKIPTWLYFYEHYSEPTLIPTFLDDDNYFSLY-KIDKDVFNDDL-LIKIKSRVLRQMXX 405 -
             +P + + Y
                           P + +D+Y+ + K D D+ L I+IK ++
Sbjct: 258 QNALPIGIP--KRYKGK---PKEIKEDHYYIYHIKADFDLKRGYLPTIQIKKKLDALRIG 312
Query: 406 XXXXXXXXXXXXXXXXXIRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQG 465
                                            + EF
Sbjct: 313 VRTSDYVTTSKNEVIDLYLTNFDLDLFLKHYDATIMYVETLE-FQTESDLFDDYI----- 366
```

```
Query: 466 KLKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYGIPALR--SHFNLFRLDDN 523
              + Y Y E+ S E +K++LN LYG + S L LDD
Sbjet: 367 -----TTYRYK-----KENAQSPAEKQKAKIMLNSLYGKFGAKIISVKKLAYLDDK 412
Query: 524 NELYNIINGYKNTERNIL-----FSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDS 577
            L +KN + + + FVTS + + ++ Q E DNF+Y DTDS
Sbjct: 413 GILR-----FKNDDEEEVQPVYAPVALFVTSIARHFIISNAQ-----ENYDNFLYADTDS 462
Query: 578 LYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAYEVNGKIKIASAGIPKN 637
L++ +L+ DP GKW E + K L K Y E+ + + K
Sbjct: 463 LHLFHSDSLVLD---IDPSEFGKWAHEGRAV-KAKYLRSKLYIEELIQEDGTTHLDV-KG 517
Query: 638 AFDTSVDFETFVREQFFDGAILENNKSIYNEQGTISIYPSKTEI 681
          A T E EFGA E ++ +G IY + +I
Sbjct: 518 AGMTPEIKEKITFENFVIGATFEGKRASKQIKGGTLIYETTFKI 561
>gi|1429230|emb|CAA67649| (X99260) DNA polymerase (Bacteriophage
          B1031
          Length = 572
 Score = 49.2 bits (115), Expect = 1e-04
 Identities = 93/422 (22%), Positives = 155/422 (36%), Gaps = 88/422 (20%)
Query: 229 KLTPEQLTYIHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTR-----FQ 283
           ++TPE+ YI ND+ I+ DI +++T + ++ + T+
Sbjct: 154 EITPEEYEYIKNDIEIIARA----LDIQFKQGLDRMTAGSDSLKGFKDILSTKKFNKVFP 209
Query: 284 LLNQYQDIKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYP 343
Query: 344 YVMYHEKIPTWLYFYEHYSEPTLIPTFLDDDNYFSLYKIDKDVFNDDLLIKIKSRVLRQM 403
MY +P Y P + + D + LY I + F +L K + +
Sbjct: 253 SQMYSRPLP-------YGAPIVFQGKYEKDEQYPLY-IQRIRFEFEL----KEGYIPTI 299
Query: 404 XXXXXXXXXXXXXXXXXXXXIRMIQ-DITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
                               ++ +T +D I+ + +Y EY
Sbjct: 300 QIKKNPFFKGNEYLKNSGAEPVELYLTNVDLELIQEH-YEMYNVEYIDGFK-----FRE 352
Query: 463 TQGKLKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPAL 511
G K I+ + H + L+K++ + LYG +P L
Sbjct: 353 KTGLFKEFIDKWTYVKTH-----EKGAKKQLAKLMFDSLYGKFASNPDVTGKVPYL 403
Query: 512 RSHFNL-FRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNF 570
           + +L FR+ D YK+ + F+T+ + + Q
Sbjct: 404 KEDGSLGFRVGDEE------YKDPVYTPM-GVFITAWARFTTITAAQACY-----DRI 449
Query: 571 IYCDTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKK-----YAYEVNG 625
           IYCDTDS+++ P + + DP LG W E+ + L K YA EV+G
Sbjct: 450 IYCDTDSIHLTGTEVPEIIKDIVDPKKLGYWAHES-TFKRAKYLRQKTYIQDIYAKEVDG 508
Query: 626 KI 627
Sbjct: 509 KL 510
 gi|1572479|emb|CAA65712| (X96987) DNA polymerase (Bacteriophage
           GA-1)
           Length = 578
 Score = 46.1 bits (107), Expect = 0.001
 Identities = 80/376 (21%), Positives = 146/376 (38%), Gaps = 54/376 (14%)
 Query: 234 QLTYIHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTRFQLLNQYQDIKI 293
++ Y+ +D++I+ + +F N D+ +T + + +Y EM + +Y +
Sbjct: 162 EIEYLKHDLLIVALA---LRSMFDN-DFTSMTVGSDALNTY--KEMLGVKQWEKYFPVL- 214
Query: 294 SYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEKIPT 353
+ I+ Y+GG N KY + + D+NS YP +M ++ +P
Sbjct: 215 -----SLKVNSEIRKAYKGGFTWVNPKYQGETVYGGMV-FDVNSMYPAMMKNKLLP- 264
 Query: 354 WLYFYEHYSEPTLIPTFLDDDNYFSLYKIDKDVFNDDLLIKIKSRVLRQMXXXXXXXXX 413
                              + + LY F + KI ++
                  Y EP +
```

```
Sbjct: 265 -----YGEPVMFKGEYKKNVEYPLYIQQVRCFFELKKDKIPCIQIKGNARFGQNEYLS 317
Query: 414 XXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLKNKINM 473

L +T +D I+ + + I+B E+ +F+ + I

Sbjct: 318 TSGDEYVDLY----VTNVDWELIKKH-YDIFEEFIGG--FMFKGF-------IGF 359
Query: 474 TSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYGIPALRSHFN--LFRLDDNNELYNIIN 531
                  + N S E+ + +K++LN LYG A + LD+N L
Sbjct: 360 FDEYIDRFMEIKNSPDSSAEQSLQAKIMLNSLYGKFATNPDITGKVPYLDENGVLKFRKG 419
Query: 532 GYKNTERNILFST---FVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYMKSVVKPLL 588
              K ER+ +++ F+T+ + N+L Q L
                                                     FIY DTDS++++ + +
Sbjct: 420 ELK--ERDPVYTPMGCFITAYARENILSNAQKLYP----RFIYADTDSIHVEGLGEVDA 472
Query: 589 NPSLFDPIALGKWDIE 604
               + DP LG WD E
Sbjct: 473 IKDVIDPKKLGYWDHE 488
>gi|118851|sp|P06950|DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2)
            >gi|75812|pir||ERBP2Z DNA-directed DNA polymerase (EC
            2.7.7.7) - phage PZA >gi|216051 (M11813) gene 2 product
            [Bacteriophage PZA] >gi|224741|prf||1112171E ORF 2
            [Bacteriophage PZA]
            Length = 572
 Score = 45.3 bits (105), Expect = 0.002
 Identities = 98/461 (21%), Positives = 166/461 (35%), Gaps = 110/461 (23%)
Query: 198 QLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYIHNDVIILGMCHIHYSDIFP 257
++ DF T+ D D + Y ++TP++ YI ND+ I+ + I
Sbjct: 129 KIAKDFKLTVLKGDIDYHKERPVGY----EITPDEYAYIKNDIQIIAEALL---IQF 178
Query: 258 NFDYNKLTFSLNIMESYLNNEMTR-----FQLLNQYQDIKISYTHYHFHDMNFYDYIKSF 312 +++T + ++ + + + T+ F L+ D ++ Y
Sbjct: 179 KQGLDRMTAGSDDLKGFKDIITTKKFKKVFPTLSLGLDKEVRYA----- 222
Query: 313 YRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEKIPTWLYFYEHYSEPTLIPT--F 370
YRGG N ++ K I E D+NS YP MY +P Y EP +
Sbjct: 223 YRGGFTWLNDRFKEKEIGEGMV-FDVNSLYPAQMYSRLLP------YGEPIVFEGKYV 273
Query: 371 LDDDNYFSLYKID-----KDVFNDDLLIKIKSRVLRQMXXXXXXXXXXXXXXXXXXXXXXIRMI 425
D+D + I K+ + + IK +SR +
Sbjct: 274 WDEDYPLHIQHIRCEFELKEGYIPTIQIK-RSRFYKGNEYLKSSGGEIADLW----- 324
Query: 426 QDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLKNKINMTSPYDYHITDDI 485
++ +D + + + +Y EY F T G K+ I+ + I
Sbjct: 325 --VSNVD-LELMKEHYDLYNVEYISGLK-----FKATTGLFKDFIDKWTHIKTTSEGAI 375
Query: 486 NEHPYSNEEVMLSKVVLNGLYG------IPALRSHFNL-FRLDDNNELYNIINGY 533
+ L+K++LN LYG +P L+ + L FRL G
 Sbjct: 376 KQ------LAKLMLNSLYGKFASNPDVIGKVPYLKENGALGFRL-----
 Query: 534 KNTERNIL--FSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYMKSVVKPLLNPS 591
 + T+ + F+T+ + Y + Q D IYCDTDS+++ P +
Sbjct: 416 EETKDPVYTPMGVFITAWARYTTITAAQACF----DRIIYCDTDSIHLTGTEIPDVIKD 470
 Query: 592 LFDPIALGKWDIENEQIDKMFVLNHKKYAY-----EVNGKI 627
             + DP LG W E+ + L K Y EV+GK+
 Sbjct: 471 IVDPKKLGYWAHES-TFKRAKYLRQKTYIQDIYMKEVDGKL 510
 >gi|2435429 (AF012250) unassigned reading frame (possible DNA
             polymerase) [Physarum polycephalum]
             Length = 544
  Score = 44.9 bits (104), Expect = 0.002
  Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)
 Query: 179 TSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYI 238
             T + LKLD + TQ F NM Y + CF L P++
 Sbjct: 62 TQLFNLLKSLQDSSFYTFKQ------FTYQNIM----YSLEISCF--LYPKKKILI 105
 Query: 239 HNDVIILGMCHIHYSDIFPNFD-----YNKL--TFSLNIMESY-LNNEMTRFQLLNQYQD 290
D+ +I Y+D+ ++ YN++ +++NI Y L+ ++ +
 Sbjct: 106 -KDLYNFFSENIIYNDVVKDYKLLAILYNEIQTAYNININRKYILSTASLSLRIFKKSFP 164
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Query: 291 IKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEK 350
                + D + +YI+ Y GG N I + + + D+NS YPY+M EK
Sbjct: 165 EKYRLIPHLTRDED--NYIRKSYIGGRNE----IFEHVAQRNYFYDVNSLYPYIMKKEK 217
Query: 351 IPTWLYFYEHYSEPTLIPTPLOD-DNYFS----LYKIDKOVFNDDLL---IKIKSRVLRQ 402
          +P + Y + + F + +N+F L I+K N +L + IK+ V
Sbjct: 218 MPIGI---PEYRDKEYMKKFEKNIENFFGFIDVLITIEKTNNNIPVLPYRMGIKNNV-EV 273
Query: 403 MXXXXXXXXXXXXXXXXXXXXXXIRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
                             L + Q I+ IY + ++++F+ Y
Sbjct: 274 GIIYAKGTLRGIYFSEBIKLALKQGYKIIE-----IYSAYEYKEKEVVFEEYVEQ 323
Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPALRS 513
                                           L K +LN LYG
              + LKK D + D
Sbjct: 324 MYNRRLKAK------DPALKD-------LYKKLLNTLYGRFGLVYEQIDIISP 363
Query: 514 HFNLFRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESBIDDNFIYC 573
            L + DN + + + + + + + + + + F Y T
Sbjct: 364 EKEL--ITDNTYISHDTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNYNLHVIYI 421
Query: 574 DTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAY-EVNGKIKIASA 632
          DTD L++K+ P+ + +L +GK+ +E+ + F+ N K Y Y +N I
Sbjct: 422 DTDGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 477
Query: 633 GIPK----NAFDTSVDFETFVR----EQFFDGAIIENNKSIYNEQGT-----ISIYPSK 678
GIP N D + + +F +I NN Y+ Q + I Y +
Sbjct: 478 GIPLQKPIFNIHDIITQHKKILNITLGHHYFTFSIRLNNNQTYSFQASRKRKLIPNYKTT 537
Query: 679 TEIVC 683
            I+C
Sbjct: 538 PWIIC 542
>gi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum
          polycephalum) >gi | 509721 | dbj | BAA06121.1 | (D29637) DNA
          polymerase [Physarum polycephalum]
          Length = 547
 Score = 44.9 bits (104), Expect = 0.002
 Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)
Query: 179 TSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYI 238
          T + L K L D + T Q F N M Y + CF L P++
Sbjct: 65 TQLFNLLKSLQDSSFYTFKQ------FTYQNIM-----YSLEISCF--LYPKKKILI 108
Query: 239 HNDVIILGMCHIHYSDIFPNFD----YNKL--TFSLNIMESY-LNNEMTRFQLLNQYQD 290
D+ +I Y+D+ ++ YN++ +++NI Y L+ ++ +
Sbjct: 109 -KDLYNFFSENIIYNDVVKDYKLLAILYNEIQTAYNININRKYILSTASLSLRIFKKSFP 167
Query: 291 IKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEK 350
K + D + +YI+ Y GG N I + + + D+NS YPY+M EK
Sbjct: 168 EKYRLIPHLTRDED--NYIRKSYIGGRNE----IFEHVAQRNYFYDVNSLYPYIMKKEK 220
Query: 351 IPTWLYFYEHYSEPTLIPTFLDD-DNYFS----LYKIDKDVFNDDLL---IKIKSRVLRQ 402 +P + Y + + F + +N+F L I+K N +L + IK+ V
Sbjct: 221 MPIGI---PEYRDKEYMKKFEKNIENFFGFIDVLITIEKTNNNIPVLPYRMGIKNNV-EV 276
Query: 403 MXXXXXXXXXXXXXXXXXXXXIRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
                                            IY + ++++F+ Y +
                             L + Q I+
Sbjct: 277 GIIYAKGTLRGIYFSEEIKLALKQGYKIIE------IYSAYEYKEKEVVFEEYVEQ 326
Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG------IPALRS 513
             + LK K D + D L K +LN LYG
Sbjct: 327 MYNRRLKAK------DPALKD-------LYKKLLNTLYGRFGLVYEQIDIISP 366
Query: 514 HFNLFRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYC 573
             L + DN + + + + + + + + + + F Y T + + IY
Sbjct: 367 EKEL--ITONTYISHOTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNYNLHVIYI 424
Query: 574 DTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAY-EVNGKIKIASA 632
           DTD L++K+ P+ + +L +GK+ +E+ + F+ N K Y Y +N I
Sbjct: 425 DTDGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 480
Query: 633 GIPK-----NAFDTSVDFETFVR----EQFFDGAIIENNKSIYNEQGT-----ISIYPSK 678
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+F +I NN . Y+ Q +
                                                           I Y +
                 N D
         GTP
Sbjct: 481 GIPLQKPIFNIHDIITQHKKILNITLGHHYFTFSIRLMNNQTYSFQASRKRKLIPNYKTT 540
Ouery: 679 TEIVC 683
Sbict: 541 PWIIC 545
>gi|4877819|gb|AAD31446.1| (AF133505) DNA polymerase [Neurospora
          crassal
          Length = 1035
Score = 44.1 bits (102), Expect = 0.004
Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)
Query: 521 DDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYM 580
          + N EL + ++G K+ 'I ++ + + ++ ++++ S Y DTDS+++
Sbjct: 817 EKNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA-----YTDTDSIFV 870
Query: 581 KSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAYEVNGKIKIASAGIPKNAFD 640
           KPL + + + K + + I + ++ K Y + GK++I GI KN +
Sbjct: 871 E---KPLDSAFIGEGCGKFKABYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 927
Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
          T+ + + E ++G + + E GT+++ K ++ G YD+
Sbjct: 928 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 977
>gi|461962|sp|P33537|DPOM_NEUCR PROBABLE DNA POLYMERASE
          >gi|283351|pir||S26985 probable DNA-directed DNA
          polymerase (EC 2.7.7.7) - Neurospora crassa
mitochondrion plasmid maranhar (SGC3)
          >gi|578156|emb|CAA39046| (X55361) putative DNA
          polymerase [Neurospora crassa]
          Length = 1021
Score = 44.1 bits (102), Expect = 0.004
Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)
Query: 521 DDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYM 580
          + N EL + ++G K+ I ++ + + ++ ++++ S Y DTDS+++
Sbjct: 815 EKNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA-----YTDTDSIFV 868
Query: 581 KSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAYEVNGKIKIASAGIPKNAFD 640
            KPL + + + K + + I + ++ K Y + GK++I GI KN
Sbjct: 869 E---KPLDSAFIGEGCGKFKAEYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 925
Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
          T+ + + E ++G + + E GT+++ K ++ G YD+
Sbjct: 926 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 975
>gi|2499511|sp|Q12471|6P22_YEAST 6-PHOSPHOFRUCTO-2-KINASE 2
          (PHOSPHOFRUCTOKINASE 2 II) (6PF-2-K 2)
          >gi|2131162|pir||S61066 6-phosphofructo-2-kinase (EC
          2.7.1.105) - yeast (Saccharomyces cerevisiae)
          >gi|2131163|pir||S71026 6-phosphofructo-2-kinase (EC
          2.7.1.105) - yeast (Saccharomyces cerevisiae)
          >gi|1420028|emb|CAA99157| (274878) ORF YOL136c
          [Saccharomyces cerevisiae] >gi|1628439|emb|CAA64733|
          (X95465) 6-phosphofructo-2-kinase [Saccharomyces
          cerevisiael
          Length = 397
Score = 40.6 bits (93), Expect = 0.041
Identities = 48/208 (23%), Positives = 92/208 (44%), Gaps = 29/208 (13%)
                                                                          Query: 175 MKTNTSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQ 234
          ++ S AT+ K LL L+ + + FN K+ND ++ +A++T ++
Sbjct: 139 IRRQISCATISKPLL----LSNTSSEDLFN----PKNNDKKET------YARITLQK 181
Query: 235 LTY-IHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTRFQLLN---QYQD 290
          L + I+ND +G+ S I + F + S+ +E++ F L+ Q
Sbjct: 182 LFHEINNDECDVGIFDATNSTI-----ERRRFIFEEVCSFNTDELSSFNLVPIILQVSC 235
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Query: 291 IKISYTHYHFHDMNFY-DYIKSFYRGGLNMYNTKYINKLIDEPCFSID-INSSYPYVMYH 348
              S+ Y+ H+ +F DY+ Y + + + + FS+D N + Y+ H
Sbjct: 236 FNRSFIKYNIHNKSFNEDYLDKPYELAIKDFAKRLKHYYSQFTPFSLDEFNQIHRYISQH 295
Query: 349 EKIPTWLYFYEHYSEPTLIPTFLDDDNY 376
            E+I T L+F+ + + P L+ +Y
Sbjct: 296 EEIDTSLFFFNVINAGVVEPHSLNQSHY 323
>gi|2258375|gb|AAD11909.1| (AP007261) transcription initiation
            factor sigma [Reclinomonas americana]
           Length = 532
 Score = 39.9 bits (91), Expect = 0.070
 Identities = 49/205 (23%), Positives = 84/205 (40%), Gaps = 14/205 (6%)
Query: 100 NHFLLKDTMRYFDNITRENIYLKSAEENEHTLKMKEATILAKNQNVIL---EKRVKSSIN 156
N++ + F + ++IY+ + +KE L K NVI+ K +K N
Sbjct: 177 NYLVKNSYLNLFKTVPHDSIYMNYSYLQTPLNILKEYLQLIKIINVIILQINKNIKKKNN 236
Query: 157 LDLTMFLNGFKFNIIDNFM---KTNTSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDND 213
           L++++FL F + N++ K + + + K L Y+T L T Y
Sbjct: 237 LNISLFLYKFYQELKWNYIFINKISRNTQKINIKTLKNSYITFYNLITFIQYYTTKKQRL 296
Query: 214 MNDSEAYDYAVKCFAK--LTPEQLTYIHNDVIILGMCHIHYSDIFPNFDYN-KLTFSLNI 270
D +K F K P+ +N +I G+ HI+ + N K+T I
Sbjct: 297 KKDIFYKQIFIKTFLKQHKIPKINKIKNNSLIKYGLTHIYDMILISILRENIKVTLKNRI 356
Query: 271 MESYLNNEMTRFQLLNQYQDIKISY 295
+ +Y+ T + QY +KI Y
Sbjct: 357 IFNYMPYITT---ISKQY--VKIGY 376
>gi|15734|emb|CAA37450| (X53370) DNA polymerase (AA 1-575)
           [Bacteriophage phi-29]
           Length = 575
 Score = 39.5 bits (90), Expect = 0.092
 Identities = 41/150 (27%), Positives = 64/150 (42%), Gaps = 36/150 (24%)
Query: 497 LSKVVLNGLYG-------IPALRSHFNL-FRLDDNNELYNIINGYKNTERNIL--F 542 L+K++LN LYG +P L+ + L FRL G + T+ +
Sbjct: 381 LAKLMLNSLYGKFASNPDVTGKVPYLKENGALGFRL---
Query: 543 STFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYMKSVVKPLLNPSLFDPIALGKWD 602
F+T+ + Y + Q D IYCDTDS+++ P + + DP LG W Sbjct: 430 GVFITAWARYTTITAAQACY----DRIIYCDTDSIHLTGTEIPDVIKDIVDPKKLGYWA 484
Query: 603 IENEQIDKMFVLNHKKYAY----EVNGKI 627
                ++ L K Y
                                EV+GK+
            E+
Sbjct: 485 HES-TFKRVKYLRQKTYIQDIYMKEVDGKL 513
Query= pt | 110872 44AHJDORF002 Phage 44AHJD ORF | 3789-5732 | 3 1
         (647 letters)
>gi|135273|sp|P27622|TAGC BACSU TEICHOIC ACID BIOSYNTHESIS PROTEIN C
           >gi|478126|pir||D49757 techoic acid biosynthesis protein
           tagC - Bacillus subtilis (strain 168) >gi|143727
           (M57497) putative (Bacillus subtilis)
           >gi|2636103|emb|CAB15594.1| (Z99122) alternate gene
           name: dinC [Bacillus subtilis]
           Length = 442
 Score = 112 bits (278), Expect = 7e-24
 Identities = 91/314 (28%), Positives = 147/314 (45%), Gaps = 58/314 (18%)
Query: 152 FELNELEPKFVMGFGGIRNAVNQSINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
                              V OS N D++ + +Y+TO S
           F+ + PK
                                                           + + I +L+ G
Sbjct: 7 FDFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLGKDNTQSYRITRLSLEG 66
Query: 208 DLISSMRIVQGGHGTTIGLERQSNGEMKIWLHHD-----GVAKLLQVAYKDNYVLDLEEA 262
             + SM + GGHGT IG+E + NG + IW +D
Sbjct: 67 LQLDSMLLKHGGHGTNIGIENR-NGTIYIWSLYDKPNETDKSELVCFPYKAGATLD-ENS 124
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Query: 263 KGLTDYTPQSLLNKHTFTPLIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDN 322
K L ++ H TP +D N +L +R + D KN+ N ++ +TI N
Sbjct: 125 KELQRFSNMPF--DHRVTPALDMKNRQLAIR------QYDTKNN--NNKQWVTIFN 170
Query: 323 SE----NNDN------RWMQGIAVDGDDLYWLSGNSSVNSHVQIGKYSLTTGQKI 367
            + N +N ++QG +D LYW +G+++ S+ +
Sbjct: 171 LDDAIANKNNPLYTINIPDELHYLQGFFLDDGYLYWYTGDTNSKSYPNL-----ITV 222
Query: 368 YDYPFKLSYQDGINFPRD-----NFKEPEGICIYTNPKTKRKSLLLAMTNGGGGKRFH 420
                                  NF+EPEGIC+YTNP+T KSL++ +T+G G R
           +D K+ Q I +D
Sbjct: 223 FDSDNKIVLQKEITVGKDLSTRYENNFREPEGICMYTNPETGAKSLMVGITSGKEGNRIS 282
Query: 421 NLYGFFQLGEYEHF 434
           +Y + YE+F
Sbjct: 283 RIYAYH---SYENF 293
>gi|142847 (M64050) DNase inhibitor (Bacillus subtilis)
           Length = 125
 Score = 51.9 bits (122), Expect = 1e-05
 Identities = 35/116 (30%), Positives = 55/116 (47%), Gaps = 10/116 (8%)
Query: 152 FELNELEPKFVMGFGGIRNAVNQSINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
                              V QS N D++ + +Y+TQ S
                                                          + + I +L+ G
Sbjct: 7 FDFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLGKDNTQSYRITRLSLEG 66
Query: 208 DLISSMRIVQGGHGTTIGLERQSNGEMKIWLHHD-----GVAKLLQVAYKDNYVLD 258
             + SM + GGHGT IG+E + NG + IW +D ++L+ YK LD
Sbjct: 67 LQLDSMLLKHGGHGTNIGMENR-NGTIYIWSLYDKPNETDKSELVCFPYKAGATLD 121
>qi|4038407 (AF103943) factor C protein precursor (Streptomyces
          griseus]
           Length = 324
 Score = 39.1 bits (89), Expect = 0.10
 Identities = 61/269 (22%), Positives = 102/269 (37%), Gaps = 33/269 (12%)
Query: 172 VNQSINIDKETNHMYSTQSDSQKPEG---FWINKLTPSGDLISSMRIVQGGHGTTIGLER 228
           V QS D ++ Q S P+
                                       I +L SG+ + M ++ GHG +IG +
Sbjct: 66 VQQSFTFDIVNRRLFVAQLKSGSPDDSGDLCITQLDFSGNKLGHMYLLGFGHGVSIGAQ- 124
Query: 229 QSNGEMKIWLHHDGVAKLLQVAYKDNYVLDLEEAKGLTDYTPQSLLNKHTFTP----- 281
+ +W D + + + + G T S L KH P
Sbjct: 125 PVGADTYLWTEVD-----VNSNARGTRLARFKWNNGATLSRTSSALAKHQPVPGATEMTC 179
Query: 282 LIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDNSENNDNRWMQGIAVDGDDL 341
ID N+++ +R+ + + + +V + V + D QG A+ G + Sbjct: 180 AIDPVNNRMAIRYLTASGRRYGIYNVADIAAGVYDKPLSDVPHPTGLGTFQGYALYGSYV 239
Query: 342 YWLSGN-----SSVNSHVQIGKYSLTTGQKIYDYPFKLSYQDGINFPRDNFKEPEGIC 394
           Y L+GN + NS+V + TG + + + G
Sbjct: 240 YQLTGNPYGPDNPNPGNSYVS--SVDVNTGALVQ----RAFTRAGSTL---TFREPEGMG 290
Query: 395 IYTNPKTKRKSLLLAMTNGGGGKRFHNLY 423
          IY + + L L +G G R NL+
Sbjct: 291 IYRTAAGEVR-LFLGFASGVAGDRRSNLF 318
Query= pt|110873 44AHJDORF003 Phage 44AHJD ORF |6626-8389|2 1
         (587 letters)
>gi|138123|sp|P04331|VG9 BPPH2 TAIL PROTEIN (LATE PROTEIN GP9)
           >gi|75850|pir||WMBPT9 gene 9 protein - phage phi-29
           >gi|215327 (M14782) tail protein [Bacteriophage phi-29]
           >gi|225364|prf||1301270D gene 9 [Bacillus sp.]
                                                                                Length = 599
 Score = 92.4 bits (226), Expect = 8e-18
 Identities = 126/618 (20%), Positives = 251/618 (40%), Gaps = 71/618 (11%)
Query: 5 TNFKFFYNTPFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPY-NFIRDRMEINVD 62
TN + + PF+ DY+NT F S+ + ++F R + + SK + F ++ ++V
Sbjct: 9 TNVRILADVPFSNDYKNTRWFTSSSNQYNWF--NRKSRVYEMSKVTFMGFRENKPYVSVS 66
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Query: 63 MQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121
                  +Y+ F + D+ ++ +YAFV ++E+ N V ++F ID + T+
Sbjct: 67 LPIDKLYSASYIMFQNADYGNKWFYAFVTELEFKNSAVTYVHFEIDVLQTWMFDMKFQES 126
Query: 122 SNVNIERQHLSKRTYNYMLPMLRNNDDVLKVSNKNYVYNQMQQYLENLVLFQSSADLSKK 181
             I R+H+ K + P + D+ L ++ + +
Sbjct: 127 F---IVREHV-KLWNDDGTPTINTIDEGLSYGSEYDIVSVENHKPYDDMMFLVIISKSIM 182
Query: 182 FGT--KKEPNLDTSKGTIYDNITSPVNLYVMEYGDFINFMDKMSAYPWITQNFQK----V 235
           GT ++E L+ ++ + + P+ Y+ + + D +I N
Sbjct: 183 HGTPGEEESRLNDINASL-NGMPQPLCYYIHPF----YKDGKVPKTYIGDNNANLSPIV 236
Query: 236 QMLPKDFINTKDLEDVKTSEKITGLKTLKQGGKSKEWSLK-DLSL-----SFSNLQ 285
                    + D+ + +T LK K+ + LK D +
Sbjct: 237 NMLTNIFSQKSAVNDI-VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVD 295
Query: 286 EMMLSK------KDEFKHMIRNEYMTIEFYDWNGNTMLLDAGKISQK 326
           + + K
                      KD+ ++ Y E D+ GN M L
Sbjct: 296 TIFVKKIPDYEALEIDTGDKWGGFTKDQESKLMMYPYCVTEITDFKGNHMNLKTEYINNS 355
Query: 327 TGVKLRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQV 386
            +K++ + +G N+V DYN+ D + N+
                                                   S +N N
Sbjct: 356 K-LKIQVRGSLGVSNKVAYSVQDYNA---DSALSGGNRLTASLDSSLINNNPN------ 404
Query: 387 PILINNGILGQSQQANRQ--KNAESQLITNRIDNVLNG---SDPKSRFYDAVSVASNLSP 441 I I N L Q N+ +N +S ++ N I ++ G + A+ + A+ + AS++
Sbjct: 405 DIAILNDYLSAYLQGNKNSLENQKSSILFNGIMGMIGGGISAGASAAGGSALGMASSV-- 462
Query: 442 TALFGKFNEEYNFYKQQQAEYKDLALQPPSVTESEMGNAFQIANSINGLTMKISVPSPKE 501
          T + + QA+ D+A PP +T+ AF N G+ +
Sbjct: 463 TGMTSTAGNAVLQMQAMQAKQADIANIPPQLTKMGGNTAFDYGNGYRGVYVIKKQLKAEY 522
Query: 502 ITFLQKYYMLFGFEVNDYNSFIEPINSMTVCNYLKCTGTYTIRDIDPMLMEQLKAILESG 561
            L ++ +G++N
                            + + NY++ + DI+ +++++ I ++G
Sbjct: 523 RRSLSSFFHKYGYKINRVKK--PNLRTRKAFNYVQTKDCFISGDINNNDLQEIRTIFDNG 580
Query: 562 VRFWHNDGSGNPMLQNPL 579
          + WHD GN ++NL
Sbjct: 581 ITLWHTDNIGNYSVENEL 598
>gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9)
          >gi|75849|pir||WMBP9Z gene 9 protein - phage PZA
          >gi|216058 (M11813) tail protein [Bacteriophage PZA]
          Length = 599
 Score = 81.9 bits (199), Expect = 1e-14
 Identities = 127/618 (20%), Positives = 248/618 (39%), Gaps = 71/618 (11%)
         TNFKFFYNTPFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPYNFIRDRME-INVD 62
          TN + + PF+ DY+NT F S+ + ++F + + SK + R+
Sbjct: 9 TNVRILADVPFSNDYKNTRWFTSSSNQYNWF--NSKTRVYEMSKVTFQGFRENKSYISVS 66
Query: 63 MQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121
                 +Y+ F + D+ ++ +YAFV ++EY N ++F ID + T+ N+ Q
Sbjct: 67 LRLDLLYNASYIMFQNADYGNKWFYAFVTELEYKNVGTTYVHFEIDVLQTW-MFNIKFQE 125
Query: 122 SNVNIERQHLSKRTYNYMLPMLRNNDDVLKVSNKNYVYN--QMQQYLENLVLFQSSADLS 179
          S I R+H+ K + P + D+ L ++ + + + + + L S +
Sbjct: 126 SF--IVREHV-KLWNDDGTPTINTIDEGLNYGSEYDIVSVENHRPYDDMMFLVVISKSIM 182
Query: 180 KKFGTKKEPNLDTSKGTIYDNITSPVNLYVMEY------GD-----FINFMDK 221
              + E L+ ++ + P+ Y+ + GD
Sbjct: 183 HGTAGEAESRLNDINASL-NGMPQPLCYYIHPFYKDGKVPKTFIGDNNANLSPIVNMLTN 241
Query: 222 MSAYPWITQNFQKVQMLPKDFINTK------DLEDVKTSEKITGLKTLKQGGKSKEWS 273
                N VM D+I K +L+ K + G+ KG
Sbjct: 242 IFSQKSAVNNI--VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVDTIFV 299
Query: 274 LKDL---SLSFSNLQEMMLSKKDEFKHMIRNEYMTIEFYDWNGNTMLLDAGKISQKTGVK 330
               +L + KD+ ++ Y E D+ GN M L I
Sbjct: 300 KKIPDYETLEIDTGDKWGGFTKDQESKLMMYPYCVTEVTDFKGNHMNLKTEYIDNNK-LK 358
Query: 331 LRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILI 390
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++ + +G N+V DYN+ + L+ + L+T++ N+ + I+

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292
Sbict: 359 IOVRGSLGVSNKVAYSIQDYNAGGS----LSGGDRLTAS----LDTSLINNNPNDIAII- 409
Query: 391 NNGILGQSQQANRQ--KNAESQLITNRIDNVLNGSDPKSRFYDAVSVASNLSP----- 441
N L Q N+ +N +S ++ N I +L G A + A SP
Sbjct: 410 -NDYLSAYLQGNKNSLENQKSSILFNGIVGMLGGG-----VSAGASAVGRSPFGLASSV 462
Query: 442 TALFGKFNEEYNFYKQQQAEYKDLALQPPSVTESEMGNAFQIANSINGLTMKISVPSPKE 501
                       + QA+ D+A PP +T+ AF N G+ +
Sbjct: 463 TGMTSTAGNAVLDMQALQAKQADIANIPPQLTKMGGNTAFDYGNGYRGVYVIKKQLKAEY 522
Query: 502 ITFLQKYYMLFGFEVNDYNSFIEPINSMTVCNYLKCTGTYTIRDIDPMLMEQLKAILESG 561
             L ++ +G+++N + + NY++ + DI+ +++++ I ++G
Sbjct: 523 RRSLSSFFHKYGYKINRVKK--PNLRTRKAYNYIQTKDCFISGDINNNDLQEIRTIFDNG 580
Query: 562 VRFWHNDGSGNPMLQNPL 579
           + WH D GN ++N L
Sbjct: 581 ITLWHTDDIGNYSVENEL 598
>gi|1429238|emb|CAA67657| (X99260) tail protein [Bacteriophage B103]
          Length = 598
 Score = 77.6 bits (188), Expect = 2e-13
 Identities = 130/623 (20%), Positives = 240/623 (37%), Gaps = 86/623 (13%)
Query: 5 TNFKFFYNTPFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPYNFI---RDRMEIN 60
          T+ + F N PF+ DY++T F + + YF + K + NF+
Sbjct: 9 TDVRIFSNVPFSNDYKSTRWFTNADAQYSYF----NAKPRVHVINECNFVGLKEGTPHIR 64
Query: 61 VDMQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLE 119
                   YM F + + ++ +Y FV ++EYVN V +YF ID I T+
          V+ + D
Sbjct: 65 VNKRIDDLYNACYMIFRNTQYSNKWFYCFVTRLEYVNSGVTNLYFEIDVIQTW-MFDFKF 123
Query: 120 QLSNVNIERQHLSKRTYNYMLPMLRNNDDVLKVSNKNYVYNQMQQYLENLVLFQSSADLS 179
          0 S + E O +
                           P+ D+ L + V Q
                                                         ++F
Sbjct: 124 QPSYIVREHQEMWDANNE---PLINTIDEGLNYGTEYDVVAVEQYKPYGDLMFMVCISKS 180
Query: 180 KKFGTKKEPNLDTSKGTIYDNITS---PVNLYVMEYGDFINFMDKMSAYPWITQNFQKVQ 236
K T E G I NI P++ YV + + D S P +T +VQ
Sbjct: 181 KMHATAGET---FKAGEIAANINGAPQPLSYYVHPF----YEDGSS--PKVTIGSNEVQ 230
Query: 237 ML-PKDFINTKDLEDVKTSEKITGLKT-----LKQGGKSKEWSLKDLSLSFSNL---- 284
          + P DF+ ++ ++ T + +K SL+D
Sbjct: 231 VSKPTDFLKNMFTQEHAVNNIVSLYVTDYIGLNIHYDESAKTMSLRDTMFEHAQIADDKH 290
Query: 285 ------QEMMLSKKDEFKHMIRNEYMTIEFY------DWNGNTMLLDAGK 322
                                     NE + Y D+ GN + +
                       +E + +F
Sbjct: 291 PNVNTIYLKEVKEYEEKTIDTGYKFASFANNEQSKLLMYPYCVTTITDFKGNQIDIKNEY 350
Query: 323 ISQKTGVKLRTKSIIGYHNEVRVYPVDYNS---AENDRPILAKNKEILIDTGSFLNTNIT 379
          ++ + +K++ + +G N+V DYN+
                                           D+ + A
Sbjct: 351 VNG-SNLKIQVRGSLGVSNKVTYSVQDYNADTTLSGDQNLTAS------CNTSLI 398
Query: 380 FNSFAQVPILINNGILGQSQQANRQ--KNAESQLITNRIDNVLN---GSDPKSRFYDAVS 434
               V I+ N L Q N+ +N + ++ N + ++L G+ +
Sbjct: 399 NNNPNDVAII--NDYLSAYLQGNKNSLENQKDSILFNGVMSMLGNGIGAVGSAATGSAVG 456
Query: 435 VASNLSPTALFGKFNEEYNFYKQQQAEYKDLALQPPSVTESEMGNAFQIANSINGLTMKI 494
                           + QA+ D+A PP + + A+ N G+ +
          VAS ST+
Sbjct: 457 VAS--SATGMVSSAGNAVLQIQGMQAKQADIANTPPQLVKMGGNTAYDYGNGYRGVYVIK 514
Query: 495 SVPSPKEITFLQKYYMLFGFEVNDYNSFIEPINSMTVCNYLKCTGTYTIRDIDPMLMEQL 554
                  L + +G++ N + + + NY++ I +++ ++++
Sbjct: 515 KQIKEEYRNILSDFSRKYGYKTNLVK--MPNLRTRESYNYVQTKDCNIIGNLNNEDLQKI 572
Query: 555 KAILESGVRFWHNDGSGNPMLQN 577
          + I +SG+ WH D G+ L N
Sbjct: 573 RTIFDSGITLWHADPVGDYTLNN 595
                                                                          _____
>gi|215339 (M12456) p9 tail protein (Bacteriophage phi-29)
          >gi|224163|prf||1011232C protein p9,tail {Bacteriophage
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phi-29] Length = 335

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Score = 71.0 bits (171), Expect = 2e-11
 Identities = 64/293 (21%), Positives = 123/293 (41%), Gaps = 20/293 (6%)
Query: 292 KDEFKHMIRNEYMTIEFYDWNGNTMLLDAGKISQKTGVKLRTKSIIGYHNEVRVYPVDYN 351
          KD+ ++ Y E D+ GN M L I+ +K++ +G N+V
Sbjct: 57 KDQESKLMMYPYCVTEITDFKGNHMNLKTEYINNSK-LKIQVRGSLGVSNKVAYSVQDYN 115
Query: 352 SAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILINNGILGQSQQANRQ--KNAES 409
                                                       Q N+ +N +S
          + D + N+ S +N N
                                            IINL
Sbjct: 116 A---DSALSGGNRLTASLDSSLINNNPN------DIAILNDYLSAYLQGNKNSLENQKS 165
Query: 410 QLITNRIDNVLNG---SDPKSRFYDAVSVASNLSPTALFGKFNEEYNFYKQQQAEYKDLA 466
            ++ N I ++ G + + A+ +AS++ T +
Sbict: 166 SILFNGIMGMIGGGISAGASAAGGSALGMASSV--TGMTSTAGNAVLOMOAMOAKOADIA 223
Query: 467 LQPPSVTESEMGNAFQIANSINGLTMKISVPSPKEITFLQKYYMLFGFEVNDYNSFIEPI 526
PP +T+ AP N G+ + + L ++ +G+++N +
Sbjct: 224 NIPPQLTKMGGNTAFDYGNGYRGVYVIKKQLKAEYRRSLSSFFHKYGYKINRVKK--PNL 281
Query: 527 NSMTVCNYLKCTGTYTIRDIDPMLMEQLKAILESGVRFWHNDGSGNPMLQNPL 579
              NY++ + DI+ +++++ I ++G+ WH D GN ++N L
Sbjct: 282 RTRKAFNYVQTKDCFISGDINNNDLQEIRTIFDNGITLWHTDNIGNYSVENEL 334
>gi|1181968|emb|CAA87738.1| (Z47794) tail protein [Bacteriophage
          CP-11
          Length = 230
 Score = 53.9 bits (127), Expect = 3e-06
 Identities = 29/113 (25%), Positives = 54/113 (47%), Gaps = 3/113 (2%)
Query: 1 MRKLTNFKFFYNTPF-TDYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPYNFIRDRMEI 59
          M++ T + +PF DY N I+F + + +D+F
                                                 + Y + + +
          MQESTKIWLYAKSPFKNDYANVINFETRESMEDFFTKKNPHIEIVYEYDKFQYTQRNGSI 60
Query: 60 NVDMQWHDAQGINYMTFLSDFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTY 112
           V + + + YM F+++ R YYAFV + Y+N+ +I + +D TY
Sbjct: 61 VVSGRVEKYENVTYMRPINN--GRTYYAFVFDVLYINEDATRIIYEVDVWNTY 111
>gi|1181970|emb|CAA87740.1| (Z47794) tail protein (Bacteriophage
          CP-1
          Length = 586
Score = 42.2 bits (97), Expect = 0.010
Identities = 79/381 (20%), Positives = 139/381 (35%), Gaps = 92/381 (24%)
Query: 277 LSLSFSNLQEMMLSK--KDEFK---HMIRNEYMTIEFYDWNGNTMLLDAG----KISQKT 327
L +++ +QE + S KD+ + ++ +E+ IE YD GN+ + I +
Sbjct: 187 LKIAYDQIQEGLRSYMGKDDLEIEVQLLNSEFTEIELYDIYGNSYVYQPQYLPRTIDEAH 246
Query: 328 GVKLRTKSIIGYHNEVRVYPVDYNSAEN----DRPIL------ 360
K+ +G N+V + ++YN+A N D+ IL
Sbjct: 247 KYKVIVSGSLGDSNQVHINFLEYNNANNVSYADKNILDSLESGDWAEHNPEHFKYGLMDV 306
Query: 361 -AKNKEILIDT-GSFLNTNITFNSFAQVPILINNGILGQSQQANRQKNAESQLITNRIDN 418
            K+ IL D S++ ++ Q+ N +L QS + ++ A +
Sbjct: 307 TGKSVAILNDAEASYIQSHKNQMEHTQLTFKENRDMLKQSVDLSNKQVATANSQASYNAQ 366
Query: 419 VLNGSDPKSRFYDAVSVASNLSPTALFGKF--------NEEYNFYKQQQ-- 459
             S +++ + S N++ LGF
                                                       N +YN QQ
Sbjct: 367 FAVDSANINOWTEGASGILNVAGNILTGNFGGALGGLASGGMKVFNANRDYNDKVVOOGF 426
Query: 460 ---------- 488
                                       A DL QP SV + AFQ N +
Sbjct: 427 TSENNALKSQSNALANMKSKIALDQSIRAYNATMADLQNQPISVQQIGNDLAFQSGNRLT 486
Query: 489 GLTMKISVPSPKEITFLQKYYMLFGFEVNDY-NSFIEPINSMTVCNYLKCTGTY--TIRD 545
                                                             T+R
                          +Y +G VN + N + + S NY+K
            K+S+ + +
Sbjet: 487 DVYWKVSLAQKEIMGRANEYIKCYGVLVNWFTNDALSVMRSRKRFNYIKMINVNLGTLR- 545
Query: 546 IDPMLMEQLKAILESGVRFWH 566
             M ++AI +SGVR W+
Sbjct: 546 ANOSHMNAIOAIFOSGVRIWN 566
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Query= pt | 110875 44AHJDORF005 Phage 44AHJD ORF | 12643-13890 | -1 1
         (415 letters)
>gi|3845203 (AE001399) GAF domain protein (cyclic nt signal
           transduct.) [Plasmodium falciparum]
           Length = 1245
 Score = 52.3 bits (123), Expect = 6e-06
 Identities = 59/246 (23%), Positives = 105/246 (41%), Gaps = 27/246 (10%)
Query: 174 ESIDRNHGNVDYIGFPKMFLLGNAVNFSSPILSNLNIYNLLQKHKMNTSRLYKNIFLEMR 233
           Sbjct: 854
Query: 234 RNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDPFYIKTDDKYI-- 291
                    + +N + M + N N ++N+ N+ N NGD Y
Query: 292 KVMYNVTTFMTNIIVVPYTKQYEFCTKIR-DIDNHVTYLRDDMFYKENMERYYYNPSNLH 350
           ++N ++ + + + K E K+ I + L +F+K NM + + L+
TSIFNKDLYVKHFVDIIMNKSLEEIIKMNVYISERINSL---LFHKGNM---LNDVTKLY 1018
Sbict: 965
Query: 351 FDNAYSKNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEKIYEDN----YIENTK 406
             NAY + N K I F + E K +M F+ +KIY+ N + N K
Sbjct: 1019 MSNAYGEKCFFFN-----FPQIKEIIFVNEYEKKMDMKYFKMLKKIYKYNLNKIFSNNYK 1073
Query: 407 KYLMKQ 412
Sbjct: 1074 FFIIKK 1079
>gi|3758843|emb|CAB11128.1| (Z98551) predicted using hexExon;
           MAL3P6.23 (PFC0820w), Hypothetical protein, len: 4982 aa
           [Plasmodium falciparum]
           Length = 4981
 Score = 49.2 bits (115), Expect = 5e-05
 Identities = 67/287 (23%), Positives = 110/287 (37%), Gaps = 60/287 (20%)
Query: 127 ITDLNSATDLKYHSNFLKHYPIIIYDEFLALEDDYLIDEWDKLKT----IYESIDRNHGN 182
I D+N + D+ + +++ I YD +++DK++ IY +ID++ N
Sbjct: 3619 IMDINKSKDISKNMEIVQN---IEYD-----NKYDKIRNDMDAIYMAIDKDMDN 3664
Query: 183 VDYIGFPKMFLLGNAVNFSSPILSNLNIYNL----LQKHKMNTSRLYKNIFLEMRRNDYV 238
+ I + F L N S +N YNL ++ K N R Y N F +D
Sbjct: 3665 IGIINCMRYFNLYKNYNNLSNECNNRE-YNLNELYMEDIKRNMKR-YDNNFNINHYDDNN 3722
Query: 239 NEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTDDKYIKVMYNVT 298
                               N N ++N N+ N NG F+ D
                 N+N++
Query: 299 TFMTNIIVVPYTKQYEFCTKIRDIDNHVTYLRDDMFYKENMERYYYNPSNLHFDNAYSKN 358
                     K FCTK
                                  ++F +N+E N N N Y+ N
Sbjct: 3772 ------KDLFFCTK------KNIFPCKNIETVCKNEYNKKIYNNYTCN 3807
Query: 359 YVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEK-IYEDNYIEN 404
                     + ++IK + + N E+ + EK +Y + EN
Sbict: 3808 ISVNNTLNCLNIIKELIKLNNNKKKILNYYEYHKVEKLLYYRHSFEN 3854
Score = 35.6 bits (80), Expect = 0.70
Identities = 62/290 (21%), Positives = 121/290 (41%), Gaps = 65/290 (22%)
           VKQNRLDMVRDYQNAVN--HVRKKIPDKYNQIELVDELMNDDIDYYISISNRSDGKSFNY 59
                     +N +N +V++ DK N I
                                            D++I+ SN + +SF
Sbjct: 4445 IKRNNINKSNIKRNNINKSNVKRSNTDKSNVIS------DFHIT-SNNNITRSFT- 4492
Query: 60 VSFFIYLAIKLDIKFTLLSRHYTLRDAYRDFIEEIIDENPLFKSKRVTFRSARDYLAIIY 119
Query: 120 QDKEIGVITDLNSATDLKYHSNFLKHYPIIIYDEFL----ALEDDYLIDEWDKLKTIYE 174
+ EI ITD++ +YH N+LK + +E+ + +D + DE ++T+ E
Sbjct: 4524 KKNEINNITDVDYGNKKEYHENYLKVKQNKVNEEYIEETFKSDKDCSIKDEACTIRTLSE 4583
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Query: 175 S--IDRNHGNVDYIGFPKMFLLGNAVNFSSPILSNLNIYNLLQKHKMN--TSRLYKNIFL 230
S I N N+D + + + S P N++ N ++K+ +N R+ KN
Sbjct: 4584 SCNISENISNID------MDDEDHISFPNGRNVHDNNYMKKNHVNYDKMRVGKNKIP 4634
Query: 231 EMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGD 280
D + +++ + +D M++ ++ E ++ + L + NG+
Sbjct: 4635 SFTHFDKILDEKKKK----SDKDMSSSKWLEREEHIKEIKLEKNEYMNGN 4680
Score = 34.0 bits (76), Expect = 2.0
 Identities = 47/211 (22%), Positives = 84/211 (39%), Gaps = 32/211 (15%)
Query: 210 IYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADD 269
Query: 270 NLRNHINQNGDFFYIKTD---DKYIKVMYNVTTFMTNIIVVPYTKQYEFCTKIRDIDNHV 326
           N+ N + D + D+ K MY + V
                                                          E K D+ N+
Sbjct: 965 NVTNEHGNHSDSYPYGNSLNLDRKPKNMYE-DIYKEKGFVKSDCSNIEI--KKNDMINND 1021
Query: 327 TYLRDDMFYKENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKII----KFHIKNE 382
Y +++ FY+++ Y+ + YV++ +YL +N ++ F +KN+
Sbjct: 1022 VYKKNE-FYEDSRINMIYDEDBIKTWFLIPHKYVIN---IIYLFLNILLTDESNFKLKNK 1077
Query: 383 MKKNMSEFERKEKIYEDN----YIENTKKY 408
E K IYEDN ++N KKY
Sbjct: 1078 KYGYFVNEETKGTIYEDNNGLQEILKNGKKY 1108
 Score = 33.6 bits (75), Expect = 2.7
 Identities = 42/198 (21%), Positives = 77/198 (38%), Gaps = 42/198 (21%)
Query: 222 SRLYKNIFLEMR---RNDYVNEKRNTRAF-----NSNDDAMTTGEFEFNEYNLA 267
S LY I++ + +N + K+NT + N+++D TT E + +
Sbjct: 411 SVLYSIIYMNKKYKKNFIITNKKNTNVYFENDVIQLSVENTSEDTFTTNTRESSLNSGM 470
Query: 268 DDNLRNHINQNGDFFYIKTDDKYIKVMYNVTTFMTNIIVVPYTKQYEFCTKIRDIDNHVT 327
            +++R +N D +DDK ++Y N
                                                   YTK E
Sbjct: 471 MNDMRYSVNNYADEKVYHSDDKSDHLIYKHVHDEKNKYDEMYTKTKE------ 517
Query: 328 YLRDDMFYKENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKFHIKNEMKKNM 387
+++ YK N+ + N K LD+ K I H+KN+ + N Sbjct: 518 --NENIIYKSNIVDKKTCDISSEMVNGKDK------LDVEKYIGSHVKND-ENNK 563
Query: 388 SEFERK-EKIYEDNYIEN 404
            + ++K + + + YI+N
Sbjct: 564 EKLKKKIDNVNKKEYIDN 581
>gi|3845297 (AE001421) hypothetical protein [Plasmodium falciparum]
            Length = 2380
 Score = 48.0 bits (112), Expect = 1e-04
 Identities = 87/390 (22%), Positives = 160/390 (40%), Gaps = 65/390 (16%)
Query: 20 VRKKIPDKYNQIELVDELMNDDIDYYISISNRSDGKSFNYVSFF----IYLAIKLDIKF 74
            +++K +K ++ + +N D + ++ R K+ NY++ +YL I DI
Sbjct: 1049 LQRKMMNKCSKNRNRNRYINKDSNIHLMNLIRIKFKNLNYMNMNSFEIELYLKINNDIFL 1108
Query: 75 TLLSRHYTLRDAYR-----DFIEEIIDEN-PLFKSKRVTFRSARDYLAIIYQDKEIGVI 127
                 +Y +++ Y + + + EN + +++ ++ +
Sbjct: 1109 QFNKHNYNVQNFYNFSITLINIMSKYYSENFYAYNLEKIVYKFLLNNKNFEYIEKQYSSK 1168
Query: 128 TDLNSATDLKYHSNFLKHYPIIIYDEFLA----LEDDYLIDEWDKLKTIYESIDRNHGNV 183
D+N D+ ++ +K+ II EFL L+ D I + KLKT ++
Sbjct: 1169 EDMNEL-DILVNTYDMKYDKII---EFLKNNGYLKIDRYIYFYPKLKT-------DI 1214
Query: 184 DYIGFPKMFLLGNAVNFSSPILSNLNIYNLLQKHKMNTSRLY-----KNIF--LEMRRN 235
                F ++FL N + L NI +++ K + Y K IF + M+ +
Sbjct: 1215 ILFFFKEIFLNDNILKIDRKFLKK-NITIMIEVLKEIFFKEYVKRCITKVIFFFVHMKEH 1273
Query: 236 DYVNEKR------NTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTD 287
            D+V K N+ FN+ D + N YN D+ N+ N N +Y K
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Sbjct: 1274 DHVMNKNYYNNQYVNNSNMFNTRGDHNNNNQTNDNHYNHHYDDTHNNNNNNNSKYY-KNK 1332
Query: 288 DKYIKVMYNVTTFMTNIIV---VPYTKQYEFCTKIRDIDNHVTYLRDDMFYKEN----ME 340
           +K K+MY +++ V K + K I + Y+ ++
Sbjct: 1333 NKN-KIMYEKERKSSSLFISNNVQDVKPIKHYLKYSSIYKNFIYIISEIKNFNNKITKIN 1391
Query: 341 RY-YYNPSNLHFDNAYSKNYVVDNDRYLYL 369
           RY YYN NL+ D+
                                ND YL+L
Sbjct: 1392 RYNYYNYMNLNIDDL-----NDAYLFL 1413
 Score = 32.5 bits (72), Expect = 6.0
 Identities = 46/183 (25%), Positives = 73/183 (39%), Gaps = 26/183 (14%)
Query: 225 YKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYI 284
          +KNI ++ ++N + NSN + +
                                            N N+ +N N IN +
Sbjct: 27 HKNINKNIKNKKFINIDNSNNCNNSNSNNSNSNNNNNNNNNIVRNN-NNFINADKKKNVI 85
Query: 285 KTDDKYIKVMYNVTTFMTNIIVVPYTKQYEFCTKIRDIDNHVTYLRDDMFYKENMERYYY 344
+D IK V NI Y ++ + D+ N+ + + KE ER
Sbjct: 86 LNEDDDIKNKELVDESFVNIFF--YENYFKNLFNLNDVSNNKVI--NIIEQKEGDER--- 138
Query: 345 NPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEKIYEDNYIEN 404
N N N +KN V DN +NK IKN +N++E Y N++ +
Sbjct: 139 NADN----NLKNKNIVRDN-----INK----IKN--TRNVNEILIYNNKYIINFLND 180
Query: 405 TKK 407
Sbjct: 181 TTK 183
>gi|4493936|emb|CAB38972.1| (AL034556) predicted using hexExon;
          MAL3P5.6 (PFC0600w), Hypothetical protein, len: 250 aa
          [Plasmodium falciparum]
          Length = 249
Score = 47.3 bits (110), Expect = 2e-04
Identities = 53/215 (24%), Positives = 87/215 (39%), Gaps = 30/215 (13%)
Query: 209 NIYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEF--NEYNL 266
          NIYN L++ YKN N ++ +N N+N
                                                        EFE N YN
Sbjct: 13 NIYNKLEEK-----YKNFLKLKNMNSHMGASQNMNV-NNNYTMNELEEFEKINNNYNN 64
Query: 267 ADDNLRNHINQNGDFFYIKTD-----DKYIKVMYNVTTFMTNIIVVPYTKQYEFCTKIRD 321
           ++N+ N+IN D+ IK +K ++ YN + I T +++
Sbjct: 65 NNNNINNNINNYYDYMNIKVSQSVQHNKRLQDFYNNKNSFQHYIKKLKTCRFDADDIRNL 124
Query: 322 IDNHVTYLRDDMFYK----ENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIK 376
          ++ + Y RD+ K EN + N + N+ S NY DN+ LY +N++ K
Sbict: 125 LEKRLAYERDNTLIKNIQEEENKKGIGINGNFGSESNSSSSNY--DNNYLLYRKINRLNK 182
Query: 377 FHIKNEMKKNMSEFERKEKIYEDNYIENTKKYLMK 411
                                  KKY++K
                         KI
Sbjct: 183 TNTNKSKNRSRKRKRINSKI-----DKKYIIK 209
>gi|3845165 (AE001390) hypothetical protein [Plasmodium falciparum]
          Length = 1247
Score = 45.7 bits (106), Expect = 6e-04
Identities = 52/239 (21%), Positives = 94/239 (38%), Gaps = 38/239 (15%)
Query: 206 SNLNIYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYN 265
          +N N +N ++K K R I +N + +N ++N+D
                                                             EN N
Sbjct: 474 NNTNKWNEIKKRKKKFKREKNKIINNSFQNQEAEDDKNNNNNDNHNDNHNDNNNENNNEN 533
Query: 266 LADDNLRNHINQNGDFFYI-KTDDKYIK----VMYNVTTFMTNIIVVPYTKQYEFCTKIR 320
                                        +YN T ++ YTK + +
           D+N N+ + N D I D+ Y
Sbjct: 534 NNDNNNENNNDINNIHNNDNNYYNNDNINLYNEMTKKKCMLDNSYTKYFFYIFTL- 592
Query: 321 DIDNHVTYLRDDMFYKENME------RYYYN-------PSNLHFDNAYS 356
            + + ++ + FY++N + ++YYN
Sbjct: 593 ---DMLPSIKFETFYEKNTDHKNFNENYKFYYNTDDDTDIINAIKKKNVKNKKKNGNIVI 649
Query: 357 KNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFER----KEKIYEDNYIENTKKYLMK 411
          KNY+ N+ Y YL+ N+ + I + K +E K+ I+ ++Y E K K
```

```
Sbjct: 650 KNYINHNE-YSYLEYNENKNYEINKKEKLLTENYEYDMYIKDNIHYNDYSEGDGKQTKK 707
 Score = 41.0 bits (94), Expect = 0.016
 Identities = 58/245 (23%), Positives = 96/245 (38%), Gaps = 43/245 (17%)
Query: 207 NLNIYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNL 266
                         Y F + D + + N D
Sbjct: 564 NINLYNEMTKKKCMLDNSYTKYFFYIFTLDMLPSIKFETFYEKNTDHKNFNENYKFYYNT 623
          N+N+YN + K K
Query: 267 ADD------NLRNHINQNGDFF---YIKTDDKYIKVMYNVT-TFMTNIIVVPYTKQ 312
                     N++N +NG+ YI ++ Y + YN + N
Sbjct: 624 DDDTDIINAIKKKNVKNK-KKNGNIVIKNYINHNE-YSYLEYNENKNYEINKKEKLLTEN 681
           ממ
Query: 313 YEFCTKIRDIDNHVTYLRDDMFYKENMERYYYNPSNLHFDNAYSK-----NYV--VD 362
          YE+ I+D ++ Y D + + YN +N +N Y K +Y+ VD
Sbjct: 682 YEYDMYIKDNIHYNDYSEGDGKQTKKASSFLYNNNN---NNKYKKEDNKTQIISYMDHVD 738
Query: 363 NDR-------YLYLDMNKIIKFHIK-NEM----KKNMSEFERKEKIYEDNYIENTKKY 408
N+ Y + +++ F +K N+M K+ F +E I + +EN K+
Sbjct: 739 NENGVKGLKKRNLFYNNSDQLYNFDVKDNDMIKYEKRQSKNFVEEEFINGNRKMENEDKH 798
Query: 409 LMKQY 413
           LKY
Sbict: 799 LKKHY 803
Query= pt|110877 44AHJDORF007 Phage 44AHJD ORF |2044-3027|1 1
         (327 letters)
>gi|1181960|emb|CAA87731.1| (Z47794) connector protein
           [Bacteriophage CP-1]
           Length = 337
 Score = 45.7 bits (106), Expect = 5e-04
 Identities = 44/184 (23%), Positives = 84/184 (44%), Gaps = 13/184 (7%)
Query: 127 QIHKLYDNCMSGNFVVMQNKPIQYNSDIEIIEHYTDELAEVALSRFSLIMQAKFSK--IF 184
           ++HK + + +V+N Y I +E + ++LA++ L+ L A+ + IF
Sbjct: 125 ELHKDNPDKIKRPCIVIPNNNF-YEPYIGYLELFCEKLADIELT-IQLNRNAQITPYFIF 182
Query: 185 KSEINDESINQLVSEIYNGAPFVKMSPMFNAD------DDIIDLTSNSVIPALTEMKR 236
N S+ + ++I N P V ++ + D D I + L ++
Sbjct: 183 ADNTNVLSMKNIFNKIANFEPVVYLNKQKDQDGQDSFKQLSDYIQVFRTDAPFLLDKLHD 242
Query: 237 EYQNKISELSNYLGINSLAVDKESGVSDEEAKSNRGFTTSNSNIYLKGREP-ITFLSKRY 295
               +++L ++GIN+ DK+ + EA SN G ++N + K R + ++K Y
Sbjct: 243 EKLRVMNQLLTFIGINNNPSDKKERLVVSEAISNNGVISANIEVGWKSRRKFVELINKCY 302
 Query: 296 GLDI 299
           GL+I
 Sbict: 303 GLEI 306
 >gi|1429239|emb|CAA67658| (X99260) upper collar protein
            [Bacteriophage B103]
            Length = 308
  Score = 44.9 bits (104), Expect = 8e-04
  Identities = 40/159 (25%), Positives = 73/159 (45%), Gaps = 11/159 (6%)
 Query: 150 YNSDIEI-----IEHYTDELAEVA-LSRFSLIMQAKFSKIFKSEINDESINQLVSEIYNG 203
                                                 I ++ N S+
                     +E + +LAE+ + + Q
           YN+D++
 Sbjct: 121 YNNDLKCSTLPALEMFAQDLAELKEIIAVNQNAQKTPVLIAANDNNQLSLKNIYNQYEGN 180
 Query: 204 APFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINSLAVDKESGV 262
 AP + + D+ + + V+ L K N E+ YLGI + ++K+ +
Sbjct: 181 APVIFVHESLDLDNLKVFKTDAPYVVDKLNAQKNAVWN---EVMTYLGIKNANLEKKERM 237
                                                                                  ---
 Query: 263 SDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
               E SN S+ NIYLK R E +S+ YGL++K
 Sbjct: 238 VTSEVDSNDEQIESSGNIYLKARQEACNKISELYGLNLK 276
 >gi|137915|sp|P07535|VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR
            PROTEIN) (LATE PROTEIN GP10) >gi|75851|pir||WMBP10 gene
```

```
10 protein - phage PZA >gi|216059 (M11813) upper collar
          protein (Bacteriophage PZA)
          Length = 309
Score = 43.8 bits (101), Expect = 0.002
Identities = 38/160 (23%), Positives = 75/160 (46%), Gaps = 13/160 (8%)
Query: 150 YNSDIEI-----IEHYTDELAEVALSRFSLIMQAKFSKIF--KSEINDESINQLVSEIYN 202
                   +E + ELAE+ S+ A+ + + + N S+ Q+ ++
Sbjct: 122 YNNDMSFPTTPTLELFAAELAELK-EIISVNQNAQKTPVLIRANDNNQLSLKQVYNQYEG 180
Query: 203 GAPFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINSLAVDKESG 261
           AP + ++D ++ + V+ L K N E+ +LGI + ++K+
Sbjct: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNAQKNAVWN---EMMTFLGIKNANLEKKER 237
Query: 262 VSDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
          + +E SN S+ ++LK R E +++ YGLD+K
Sbjct: 238 MVTDEVSSNDEQIESSGTVFLKSREEACEKINELYGLDVK 277
>gi|137914|sp|P04332|VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR
          PROTEIN) (LATE PROTEIN GP10) >gi | 75852 |pir | WMBPC9 gene
          10 protein - phage phi-29 >gi | 215328 (M14782) upper
          collar protein (Bacteriophage phi-29) >gi|215340
          (M12456) pl0 connector protein [Bacteriophage phi-29]
          >gi|224161|prf||1011232A protein pl0,connector
          [Bacteriophage phi-29] >gi|225365|prf||1301270E gene 10
          [Bacteriophage phi-29]
          Length = 309
Score = 41.4 bits (95), Expect = 0.009
Identities = 37/160 (23%), Positives = 75/160 (46%), Gaps = 13/160 (8%)
Query: 150 YNSDIEI-----IEHYTDELAEVALSRFSLIMQAKFSKIF--KSEINDESINQLVSEIYN 202
                   +E + ELAE+ S+ A+ + + + N S+ Q+ ++
          YN+D+
Sbict: 122 YNNDMAFPTTPTLELFAAELAELK-EIISVNQNAQKTPVLIRANDNNQLSLKQVYNQYEG 180
Query: 203 GAPFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINSLAVDKESG 261
AP + ++D ++ + V+ L K N E+ +LGI + ++K+
Sbjct: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNAQKNAVWN---EMMTFLGIKNANLEKKER 237
Query: 262 VSDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
          + +E SN S+ ++LK R E +++ YGL++K
Sbjct: 238 MVTDEVSSNDEQIESSGTVFLKSREEACEKINELYGLNVK 277
Query= pt | 110878 44AHJDORF008 Phage 44AHJD ORF | 3020-3775 | 2 1
        (251 letters)
>gi|4982468|gb|AAD30963.2| (AF118151) SNF1/AMP-activated kinase
          [Dictyostelium discoideum]
         Length = 718
Score = 52.3 bits (123), Expect = 3e-06
Identities = 28/118 (23%), Positives = 56/118 (46%), Gaps = 5/118 (4%)
Query: 121 YLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYV----SLPQSEVNIDVDN 176
          + + GF N ++ SN + +N N + N+ T N N + ++ +N + +N
Query: 177 TTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLID-NIDKAYD 233
               +NN I+N N ++N +N N N N N + + T+ + I N++ +Y+
Score = 37.5 bits (85), Expect = 0.094
Identities = 17/111 (15%), Positives = 45/111 (40%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
         +N + +N + +N N + +N++ ++ + P
Sbjct: 456 NNNNNNNNNNNNNNNNNNNNNNNNNNSSISGGTEVFSISPNLNNSYNSNSSGNSNGSNSNNNS 515
Query: 190 GKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKKILN 240
             N +N +N N N N N ID+++ + + N
Sbjct: 516 NNNTNNDNNNNNNNNNNNNNNNNNNNNNNNNNCIDSVNNSLNNENDVNN 566
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Score = 32.8 bits (73), Expect = 2.4
 Identities = 31/140 (22%), Positives = 57/140 (40%), Gaps = 14/140 (10%)
 Query: 109 LNVVYSSSEVEKYLQSQGFTEHNEDTTS---NTDETSNQNATSLDNSTGMTANRNAYVSL 165
         LN Y+S+
                          N +T + N + +N N + +N+
                    S
 Query: 166 PQSEVN--IDVDNTTLRFADNNTIDNGKTVNKSS-----NESNQNAKRNQNQKGNAK 215
            + +N DV+N+ + +NN D+G N
                                           ++ N N + N GN
 Sbjct: 554 VNNSLNNENDVNNSNINNNNNNSDDGSNNNSYEGGGDVLLLSDLNGNNQLGGNDNGNVV 613
 Query: 216 GTQFTKQYLIDNIDKAYDLR 235
              O L++++D D++
 Sbjct: 614 NLNNNFQ-LLNSLDLNSDIQ 632
 Score = 31.7 bits (70), Expect = 5.4
 Identities = 25/115 (21%), Positives = 48/115 (41%), Gaps = 10/115 (8%)
Query: 130 HNEDTTSNTDETSNQNATSLDNST---GMTAN-RNAYVSLPQSEVNIDVDNTTLRFADNN 185 +N + +N + +N N +S+ T ++ N N+Y S S N + N+ +N Sbjct: 462 NNNNNNNNNNNNNNNNNSSISGGTEVFSISPNLNNSYNS--NSSGNSNGSNSNNNSNNNT 519
Query: 186 TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKKILN 240
           DN N ++N +N N N N
                                        + ++++ D+
Score = 31.7 bits (70), Expect = 5.4
 Identities = 15/104 (14%), Positives = 43/104 (40%)
Query: 110 NVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSE 169
         N+ +++ + +N + +N N + +N+ + +
Query: 170 VNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
         +N ++ ++ ++ +N N +++ +N N N
Sbjct: 494 LNNSYNSNSSGNSNGSNSNNNSNNNTNNDNNNNNNNNNNNNNNN 537
 Score = 30.9 bits (68), Expect = 9.2
 Identities = 16/84 (19%), Positives = 34/84 (40%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
         Sbict: 455 NNNNNNNNNNNNNNNNNNNNNNNNNSSISGGTEVFSISPNLNNSYNSNSSGNSNGSNSNNN 514
Query: 190 GKTVNKSSNESNQNAKRNQNQKGN 213
              + N +N N N N
Sbjct: 515 SNNNTNNDNNNNNNNNNNNNNNNN 538
>gi|1730077|sp|P18160|KYK1_DICDI NON-RECEPTOR TYROSINE KINASE SPORE
         LYSIS A (TYROSINE-PROTEIN KINASE 1) >gi|974334 (U32174)
         non-receptor tyrosine kinase [Dictyostelium discoideum]
         Length = 1584
 Score = 46.5 bits (108), Expect = 2e-04
 Identities = 29/106 (27%), Positives = 48/106 (44%), Gaps = 4/106 (3%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNID---VDNTTLRFADN-N 185
         +NED +SN + +N N + +N+ N N + + N + ++NTT
Query: 186 TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKA 231
               N +SN +N N N N N TK+ I + D++
          +N
Score = 34.0 bits (76), Expect = 1.1
Identities = 20/117 (17%), Positives = 46/117 (39%)
Query: 87 NRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQNA 146
               G IT T + + ++ ++
                                           +N + +N + +N N
        N
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Sbjct: 415 NNNNNNIIGNGKITTTTTTSTSPSSINNNEDISSNNNNNNNNNNNNNNNNNNNNNNNNNN 474
Query: 147 TSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQN 203
           + ++++ T N N + + N + +N N ++N N
Score = 33.2 bits (74), Expect = 1.8
 Identities = 18/88 (20%), Positives = 35/88 (39%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
          +N + ++N + +N N T T + S+ +E +N +NN +N
Sbjct: 405 NNNNNSNNNNNNNNNNIIGNGKITTTTTTSTSPSSINNNEDISSNNNNNNNNNNNNNNNNN 464
Ouery: 190 GKTVNKSSNESNONAKRNONOKGNAKGT 217
             N ++N +N N+ + N T
Score = 32.5 bits (72), Expect = 3.1
 Identities = 18/94 (19%), Positives = 37/94 (39%)
Query: 120 KYLQSQGFTEHNEDTTSNTDBTSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTL 179
         K + S N + +N++ +N N ++ + +T S
                                                    N D+ +
Sbjct: 392 KNVNSTSILVPNGNNNNNNNNNNNNNNNNNIIGNGKITTTTTSTSPSSINNNEDISSNNN 451
Query: 180 RFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
            +NN +N N ++N N + + N
Score = 32.5 \text{ bits } (72), \text{ Expect = } 3.1
 Identities = 24/110 (21%), Positives = 44/110 (39%), Gaps = 10/110 (9%)
Query: 138 TDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGK----- 191
         T T++ + +S++N+ +++N N + + N + +N
                                                 +NN
Query: 192 ----TVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
             TN+SN+NNNNN+
>gi|3758855|emb|CAB11140.1| (Z98551) predicted using hexExon;
         MAL3P6.11 (PFC0760c), Hypothetical protein, len: 3395 aa
          (Plasmodium falciparum)
         Length = 3394
 Score = 46.5 bits (108), Expect = 2e-04
 Identities = 52/202 (25%), Positives = 96/202 (46%), Gaps = 32/202 (15%)
Query: 21 FNEFVNDNKLTFYDDEFQFMQKMLKFD-KDVLAIVNEKVFKGFSLKDELSDL--LFKKSF 77
F ++ ++ K T D+ M+K K D DV + NEK++ L ++L+ + + KK
Sbjct: 665 FEKYCSNIKNTLIRDD---MKKFRKPDISDVHILHNEKIYLEKLLNEKLNYIKDIEKKLD 721
Query: 78 TIHFLDRBINRQTVEAFGMQV----ITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNE 132 +H + IN+ + + +QV I V + DY + S + + K + +N
Sbjct: 722 ELHGV---INKNKEDIYILQVEKQTLIKVISSVYDYTKME-SENHIFKMNTTWNKMLNNV 777
Query: 133 DTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKT 192
           +SN D +NQN +++N+ + N+N N +++N + N +N
Sbjct: 778 HMSSNKDY-NNQNNQNIENNQNIENNQN------NQNIEN-----NQNIENNQNN 820
Query: 193 VNKSSNESNQNAKRNQNQKGNA 214
          N +N++NQN + NQN + NA
Sbjct: 821 QNNQNNQNNQNNQNNQNNQNNA 842
Score = 33.6 bits (75), Expect = 1.4
                                                                        ____
Identities = 46/221 (20%), Positives = 89/221 (39%), Gaps = 37/221 (16%)
Query: 10 DFIKSELIKKGFNEFVNDNKLTFYDDEFQFMQKMLKFDKDVLAIVNEKVFKGFSLKDELS 69
         D + K E K N + + L Y + + M+K K
                                                + V K SL
Sbjct: 367 DSLKIEYNKSKTNIQQLNEQLVNYKNFIKEMEKKYK------QLVVKNNSLFSITH 416
Query: 70 DLLFKKSFTIHFLDREINRQTVEAFGMQVITVCITH---EDYLNVVYSSSEVEKYLQSQG 126
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D + K+ I + R + + +
                                  ++ + I H +D+L+V+Y
Sbjct: 417 DFINLKNSNIIIIRRTSDMKQI----FKMYNLDIEHFNEQDHLSVIY----IYEILYNTN 468
Ouery: 127 FTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186
             +N D +N D +N N + +N+
                                      N N
                                                  N + +N +
Query: 187 IDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDN 227
          I+N + N +++ N + N N + N +++Y I+N
Sbict: 513 IENMNSGNHPNSNNLHNYRHNTNDENNLSSLKTSFRYKINN 553
 Score = 32.8 bits (73), Expect = 2.4
 Identities = 28/122 (22%), Positives = 53/122 (42%), Gaps = 2/122 (1%)
Query: 119 EKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNID-VDNT 177
Query: 178 TLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
                +N+ +NG SSN ++ N N N K N +G + + + + YD K
Sbjct: 2898 NNNDNNNDNSNNGFVCELSSNINDFNNILNVN-KDNFQGINKSNNFSTNLSEYNYDAYVK 2956
Query: 238 IL 239
Sbjct: 2957 IV 2958
 Score = 32.5 bits (72), Expect = 3.1
 Identities = 46/249 (18%), Positives = 101/249 (40%), Gaps = 31/249 (12%)
           YDFIKSELIKKGFNEFVNDNKLTFYDDEFQFMQKMLKFDKDVLAIVNEKVFKGFSLKDEL 68
Y+++K ++ N N NK B Q++ K+ + + + E K L++
Sbjct: 2150 YNYVK---VQNATNREDNKNK-----ERNLSQEIYKYINENIDLTSELEKKNDMLENYK 2200
           SDL-----LFKKSFTIHFLDREINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYL 122
Query: 69
++L ++K + I L + M+ + + N + E+ + L
Sbjct: 2201 NELKEKNEEIYKLNNDIDMLSNNCKKLKESIMMMEKYKIIMN-----NNIQEKDEIIENL 2255
Query: 123 QSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTAN----RNAYVSLPQSE----VNIDV 174 +++ +D +N + ++S M+ + N + +L +S N+D+
           +++ + +D +N
Sbjct: 2256 KNK-YNNKLDDLINNYSVVDKSIVSCFEDSNIMSPSCNDILNVFNNLSKSNKKVCTNMDI 2314
Query: 175 DNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDL 234
                                                       YL++N+
                  ++I+N +N +N +N N N N K
Sbict: 2315 CNENMDSI--SSINNVNNINNVNNINNVNNINNVKNIVDINNYLVNNLQLNKDN 2372
Query: 235 RKKILNEFD 243
              I+ +F+
Sbjct: 2373 DNIIIIKFN 2381
 Score = 32.1 bits (71), Expect = 4.1
 Identities = 20/103 (19%), Positives = 48/103 (46%), Gaps = 2/103 (1%)
Query: 115 SSEVEKYLQSQGFTEHNEDTTSNTDETSNQN--ATSLDNSTGMTANRNAYVSLPQSEVNI 172
                     EH + N D +N+N
           +++ EKY
                                            L ++ ++ + N S ++R+
Sbjct: 3264 NNDEEKYSCHDDKNEHTNNDLLNIDHDNNKNNITDELYSTYNVSVSHNKDPSNKENEIQN 3323
Query: 173 DVDNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAK 215
            + + DN ++ N ++E+++N + ++N + + K
Sbjct: 3324 LISIDSSNENDENDENDENDENDENDENDENDENDENDENDEK 3366
Score = 30.9 bits (68), Expect = 9.2
Identities = 27/118 (22%), Positives = 53/118 (44%), Gaps = 15/118 (12%)
Query: 104 THEDYLNVVYSSSEV----EKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANR 159
           T+ D LN+ + +++ E Y HN+D ++ +E QN S+D+S N
Sbjct: 3280 TNNDLLNIDHDNNKNNITDELYSTYNVSVSHNKDPSNKENEI--QNLISIDSSNENDEND 3337
Query: 160 NAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                 +++ N + D D N ++ N +E+++N + ++N N +GT
Sbict: 3338 En----DENDENDENDEN-----DENDENDENDENDENDENDENDENFDNNNEGT 3386
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>gi|585795|sp|P21538|REB1_YEAST DNA-BINDING PROTEIN REB1 (QBP)
           -gi|626139|pir||S45907 DNA-binding protein REB1 - yeast
           (Saccharomyces cerevisiae) >gi|536280|emb|CAA84992|
           (235918) ORF YBR049c [Saccharomyces cerevisiae]
           >gi|559944|emb|CAA86391| (Z46260) REB1 DNA-binding
           protein [Saccharomyces cerevisiae]
           Length = 810
 Score = 45.7 bits (106), Expect = 3e-04
 Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)
Query: 83 DREINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETS 142
           D+ N+++VE ++ + V + H+++ +++
                                               K+ + Q E + D N ++ S
          DKNANQESVEEAVLKYVGVGLDHQNHDPQLHTKDLENKHSKKQNIVESSSDVDVNNNDDS 66
Sbict: 7
Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
N+N + D+S ++A L +E + +VD+ N +D N+ +E
Sbjct: 67 NRNEDNNDDSENISA-----LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119
Query: 200 SNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
            ++N N GN F++ ++ +D D KK
Sbjct: 120 DDEN--NNNTDNGNDSNNHFSQSDIV--VDDDDDKNKK 153
>qi|172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]
          Length = 809
 Score = 45.7 bits (106), Expect = 3e-04
 Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)
Query: 83 DREINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETS 142
          D+ N+++VE ++ + V + H+++ +++ K+ + Q E + D N ++ S
          DKNANOESVEEAVLKYVGVGLDHQNHDPQLHTKDLENKHSKKQNIVESSNDVDVNNNDDS 66
Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
                                                  N +D
          N+N + D+S ++A L +E + +VD+
Sbjct: 67 NRNEDNNDDSENISA------LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119
Query: 200 SNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
++N N GN F++ ++ +D D KK
Sbjct: 120 DDEN--NNNTDNGNDSNNHFSQSDIV--VDDDDDDKNKK 153
>gi|2952545 (AF051898) coronin binding protein [Dictyostelium
          discoideuml
          Length = 560
 Score = 44.9 bits (104), Expect = 6e-04
 Identities = 26/83 (31%), Positives = 39/83 (46%), Gaps = 5/83 (6%)
Query: 131 NEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNG 190
          N + +N +N N+ S +NS +N N+ + P
                                                 N D DN T +NNT +N
Sbjct: 404 NNNNNNNIINNNNSNSNSNNNSNN-NSNNNSNRNSPNHNNNGDNDNNT----NNNTNNNN 458
Query: 191 KTVNKSSNESNQNAKRNQNQKGN 213
             N ++N +N N N N
Sbjct: 459 NNNNNNNNNNNNNNNNNNNNNN 481
 Score = 41.4 bits (95), Expect = 0.006
 Identities = 22/88 (25%), Positives = 43/88 (48%), Gaps = 6/88 (6%)
Query: 130 HNEDTTSNTDETSNQNATSLDN---STGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186
+ ++ +N++ SN N+ +N +G AN++ +P + +N + DN +NN
Sbjct: 337 NRNNSNNNSNNNSNNSNNSNNRNITNGSNANKS---NSPNNNLNTNNDNKNNNSNNNNN 393
Query: 187 IDNGKTVNKSSNESNQNAKRNQNQKGNA 214
                  S+N +N N N N+
Sbjct: 394 SNNNSNNGNSNNNNNNNIINNNNSNSNS 421
Score = 40.6 bits (93), Expect = 0.011
 Identities = 24/101 (23%), Positives = 41/101 (39%), Gaps = 2/101 (1%)
Query: 115 SSEVEKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDV 174
                       ++N +N ++ N S +N+ N N S + N +
          S+
                 L +
```

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Query: 175 DNTTLRFADN--NTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
          +N + R + N N DN N ++N +N N N N
Sbjct: 430 NNNSNRNSPNHNNNGDNDNNTNNNTNNNNNNNNNNNNNNNN 470
 Score = 40.2 bits (92), Expect = 0.014
 Identities = 21/80 (26%), Positives = 39/80 (48%), Gaps = 9/80 (11%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
         +N D +NT+ +N N + +N+ N N N + +N
                                                      +ADN+ ++
Query: 190 GKTVNKSSNESNQNAKRNQN 209
           + N +SN +N N +N+N
Sbjct: 493 SNSNNNNSNSNNNNDNKNEN 512
 Score = 39.5 bits (90), Expect = 0.024
 Identities = 26/111 (23%), Positives = 44/111 (39%), Gaps = 20/111 (18%)
Query: 112 VYSSSEVEKYLQSQ--GFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSE 169
         VY + K+ ++ G +N ++ +N++ SN N ++N
Sbict: 296 VYCTHHHTKFYETHRNGLLNNNNNSNNNSNSNSNNNNNGINNRNNSNNNSN----- 346
Query: 170 VNIDVDNTTLRFADNNTIDNGKTVNKSS-----NESNQNAKRNQNQKGNA 214
                  ++N I NG NKS+ N +N N N N+
Sbict: 347 --- NISNINISNINSNIRNITNGSNANKSNSPNNNLNTNNDNKNNNSNNNNNS 394
Score = 37.5 bits (85), Expect = 0.094
Identities = 24/96 (25%), Positives = 41/96 (42%), Gaps = 1/96 (1%)
Query: 124 SQGFTEHNEDTTSNTDETSNQNATSLDNSTGM-TANRNAYVSLPQSEVNIDVDNTTLRFA 182
         S + +N + SN + ++ N DN+T T N N
                                                 + N + +N
Sbict: 421 SNNNSNNNSNNNSNRNSPNHNNNGDNDNNTNNNTNNNNNNNNNNNNNNNNNNNNNNN 480
Query: 183 DNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQ 218
         +NN DN + +SN +N N+ N + K
Sbjct: 481 NNNYADNSNNNSSNSNNNNSNSNNNNDNKNENSDNQ 516
Score = 35.6 bits (80), Expect = 0.36
Identities = 25/99 (25%), Positives = 42/99 (42%), Gaps = 18/99 (18%)
Query: 130 HNEDTTSNTDETSNQNATSLDNST-GMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID 188
+N + SN + +N N ++ N T G AN++ + P + +N + DN +NN +
Sbjct: 339 NNSNNNSNNNSNNSNNRNITNGSNANKS---NSPNNNLNTNNDNKNNNSNNNNNN 395
Query: 189 NGKTV------NKSSNESNQNAKRNQNQKGN 213
                          N S++ SN N+ N N
Sbjct: 396 NNSNNGNSNNNNNNNIINNNNSNSNSNNNSNNNSNNNSN 434
Score = 35.2 bits (79), Expect = 0.47
Identities = 21/94 (22%), Positives = 42/94 (44%), Gaps = 5/94 (5%)
Query: 124 SQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFAD 183
Query: 184 NNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
         +N+ N + N S+N SN+N+ + N N T
Sbjct: 417 SNSNSNNNSNNNSNNNSNRNSPNHNNNGDNDNNT 450
Score = 35.2 bits (79), Expect = 0.47
Identities = 29/118 (24%), Positives = 53/118 (44%), Gaps = 12/118 (10%)
Query: 115 SSEVEKYLQS-QGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNID 173
         SS+ E ++ +GF + + T+N ++N
                                       D S+G + + + V+ P+S +N
Sbjct: 114 SSDSEADIEDDKGFQD--KPITTNNSGSNNPLKNLKDYSSGSSGSSRSGVNQPRSNINNS 171
Query: 174 VDNTTLRFADNNT------IDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQ 222
          D + + +N+
                           I + T + NQN +NQNQ N
```

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Sbjct: 172 NDKYKSKSSSSNSNSSSSGGSLISSLLTGGNTYQNQNQNQNQNQNQNNNQQQLQQQQQ 229
 Score = 34.4 bits (77), Expect = 0.81
 Identities = 24/94 (25%), Positives = 38/94 (39%), Gaps = 12/94 (12%)
 Query: 131 NEDITSNIDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNITLRFADNNTIDNG 190
            N +T +N + +N N + +N+ N N
                                                 s n
 Sbjct: 451 NNNTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNSNSN----
Query: 191 KTVNKSSNESNQNAKR-----NQNQKGNAKGTQ 218
NK+ N NQ+ R ++NQK + Q
 Sbjct: 505 NNDNKNENSDNQSVLRSNEKFTDENQKNGSDDQQ 538
 Score = 33.6 bits (75), Expect = 1.4
 Identities = 22/90 (24%), Positives = 35/90 (38%)
Query: 124 SQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFAD 183
                 N SN ++++ N
                                     N+ N N + + N + +N
Sbjct: 353 SNNSNNRNITNGSNANKSNSPNNNLNTNNDNKNNNSNNNNSNNNSNNGNSNNNNNNNNII 412
Query: 184 NNTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
           NN N + N S+N SN N+ RN
Sbjct: 413 NNNNSNSNSNNNSNNNSNRNSPNHNN 442
>gi|535260|emb|CAA82996| (230339) STARP antigen [Plasmodium
           reichenowil
           Length = 655
 Score = 44.5 bits (103), Expect = 7e-04
 Identities = 31/114 (27%), Positives = 47/114 (41%), Gaps = 14/114 (12%)
Query: 128 TEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVN-----IDVDNTTLRF 181
           T++N T TD + + +N+T A N + ++ N
                                                               D +NT +
Sbjct: 433 TDNNNTNTKATDSNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKA 492
Query: 182 ADNNTI-----DNGKTVNKSSNESNQNAKRNQNQKGNAKGT---QFTKQYLIDN 227
                     DN T K+++ +N N K N N K T
Sbjct: 493 TDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNNTNQYVFAN 546
 Score = 44.5 bits (103), Expect = 7e-04
 Identities = 30/103 (29%), Positives = 44/103 (42%), Gaps = 13/103 (12%)
Query: 128 TEHNEDTTSNTDETSNQNATSLDNS----TGMTANRNAYVSLPQSEVN----IDVDNTTL 179
           T++N T TD+++N + + DN+ T T N N
Sbjct: 401 TDNNNTDTKATDKSNNTDTKATDNNNNTDTKATDNNNTNTKATDSNNTNTKATDNNNTNT 460
Query: 180 RFADNNTI-----DNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                      DN T K+++ +N N K N N K T
           + DNN
Sbjct: 461 KATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 503
 Score = 42.6 bits (98), Expect = 0.003
 Identities = 27/96 (28%), Positives = 43/96 (44%), Gaps = 10/96 (10%)
Query: 128 TEHNEDTTSNTDETSNQNATSLD-NSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186 T++N +T + + +N N + D N+T A N + ++ N NT + DNN Sbjct: 422 TDNNNTDTKATDNNNTNTKATDSNNTNTKATDNNNTNTKATDNNN----NTNTKATDNNN 477
Query: 187 I----DNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                 DN T K+++ +N N K N N K T
Sbjct: 478 TNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 513
 Score = 41.8 bits (96), Expect = 0.005
 Identities = 35/150 (23%), Positives = 59/150 (39%), Gaps = 9/150 (6%)
Query: 85 EINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQ 144 E N+ ++ G T+ + N + E + +Q T +N TT+ + N
Sbjct: 118 ETNKTNIKLTGNNSTTINTNLTENTNA--TKKLTENVITNQILTGNNNTTTNTSSTEHNN 175
Query: 145 NATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNA 204
                                     NI + N L +N T + T + ++ +N N+
           N + NSTG T+
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Sbjct: 176 NINTNTNSTGNTSTTKKLTE-----NI-ITNQILTGNNNTTTNTSSTEHNNNINTNTNS 228
Query: 205 KRNQNQKGNAKGTQFTKQYLIDNIDKAYDL 234
            N N N
                     T + DNI+ +L
Sbjct: 229 TDNSNTNTNLTDITTTTKKWTDNINTTQNL 258
 Score = 41.8 bits (96), Expect = 0.005
 Identities = 30/101 (29%), Positives = 43/101 (41%), Gaps = 13/101 (12%)
Query: 130 HNEDTTSNTDETSNQNATSLDNS-TGMTANRNAYVSLPQSEVNIDV-----DNTTLRFA 182
          +N DT S ++ ++ AT DN+ T T N N +
                                                 N D
Sbjct: 363 NNTDTISTDNDNTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKAT 422
Query: 183 DNN-----TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
          DNN DN T K+++ +N N K N N K T
Sbjct: 423 DNNNNTDTKATDNNNTNTKATDSNNTNTKATDNNNTNTKAT 463
 Score = 40.6 bits (93), Expect = 0.011
 Identities = 31/121 (25%), Positives = 47/121 (38%), Gaps = 31/121 (25%)
Query: 128 TEHNEDTTSNTDETSNQNAT-----SLDNSTGMTANRNAYVSLPQSEVN------- 171
TEHN + +NT+ T N + T ++ +T N N + +E N
Sbjct: 171 TEHNNNINTWINSTGNTSTTKKLTENIITNQILTGNNNTTTNTSSTEHNNNINTNTNSTD 230
Query: 172 -----IDVDNTTLRFADN-----NTIDNGKTVNKSSNESNQNAKRNQNQKGNAKG 216
                  D+ TT ++ DN T N TV+ +N +N N K N N K
Sbjct: 231 NSNTNTNLTDITTTTKKWTDNINTTQNLTTSTNTTTVSTDNNNNNINTKPTDNNNTNIKS 290
Ouerv: 217 T 217
Sbjct: 291 T 291
 Score = 38.3 bits (87), Expect = 0.055
 Identities = 28/98 (28%), Positives = 41/98 (41%), Gaps = 10/98 (10%)
Query: 128 TEHNEDITSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVD-NTTLRFADNNT 186
          TEHN + +NT+ S N+ + N T +T + + N+
                                                        NTT
Sbjct: 216 TEHNNNINTNTN--STDNSNTNTNLTDITTTTKKWTDNINTTQNLTTSTNTTTVSTDNNN 273
Query: 187 -----IDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                  DN T KS++ N K N+ + K T
Sbjct: 274 NNINTKPTDNNNTNIKSTDNYNTGTKETDNKNTDIKAT 311
Score = 37.5 bits (85), Expect = 0.094
 Identities = 31/106 (29%), Positives = 45/106 (42%), Gaps = 18/106 (16%)
Query: 128 TEHNEDTTSNTDETSNQN----ATSLONSTGMTANRNAYVSLPQSEVN-----IDVDN 176
          T++N +T +T T N N AT N+T A N + ++ N
Sbjct: 390 TDNNNT--DTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKATDNNNTNTKATDSNN 447
Query: 177 TTLRFADNN-----TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
          T + DNN
                        DN T K+++ +N N K N N K T
Sbjct: 448 TNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 493
Score = 35.2 bits (79), Expect = 0.47
Identities = 24/109 (22%), Positives = 46/109 (42%), Gaps = 6/109 (5%)
Query: 128 TEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVN-----IDVDNTTLRF 181
          T++N T TD + + +N+T A N + ++ N
Sbjct: 473 TDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKA 532
Query: 182 ADMNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDK 230
                                                                           DNN N + +E+ + K N++ N++ + K + +DK
Sbjct: 533 TDNNNNTNQYVFANNYDETTSDDKLNKDSCDNSEEKENIKSMINAYLDK 581
Score = 34.4 bits (77), Expect = 0.81
Identities = 26/126 (20%), Positives = 46/126 (35%), Gaps = 7/126 (5%)
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(250 letters)

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Query: 99 ITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTAN 158
          IT T+ + ++ S + V S T +++ +N T N N ++
Sbjct: 318 ITTONTNTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVISTDNNNTDTISTDNDNTDT 377
Query: 159 RNAYVSLPQSEVNIDVDNTTLRFADNNTID-----NGKTVNKSSNESNQNAKRNQNQK 211
                   ++ + +NT + DNN D N + N +N + K N
Sbict: 378 KATDNDNTDTKATDNNNTDTKATDNNNTDTKATDNNNTDTKATDNNNTDTKATDNNN 437
Query: 212 GNAKGT 217
           NKT
Sbjct: 438 TNTKAT 443
 Score = 34.4 bits (77), Expect = 0.81
 Identities = 30/100 (30%), Positives = 44/100 (44%), Gaps = 14/100 (14%)
Query: 131 NEDTTSNTDETSNQNATSLDNS-TGMTANRNAY---VSLPQSEVNI---DVDNTTLRFAD 183
           N + T TD T N N S DNS T + + N+ +S S+ N+ D +NT
Sbict: 313 MNNITITTDNT-NTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVISTDNNNTDTISTD 371
Query: 184 NNTIDNGKTVNKSS-----NESNQNAKRNQNQKGNAKGT 217
N+ D T N ++ N +N + K N + K T
Sbjct: 372 NDNTDTKATDNDNTDTKATDNNNNTDTKATDNNNTDTKAT 411
 Score = 34.4 bits (77), Expect = 0.81
 Identities = 28/101 (27%), Positives = 41/101 (39%), Gaps = 15/101 (14%)
Query: 131 NEDTTSNTDETSNQNATSLDNSTGMTA--NRNAYVSLPQSEVNIDV-----DNTTLRFA 182
           N DT + ++ ++ AT +N+T A N N
                                                    N D
Sbjct: 374 NTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKAT 433
Query: 183 DNNTIDNGK-----TVNKSSNESNQNAKRNQNQKGNAKGT 217
          DNN NK T K+++ +N NK N NK T
Sbjct: 434 DNNN-TNTKATDSNNTNTKATDNNNTNTKATDNNNTNTKAT 473
 Score = 32.5 bits (72), Expect = 3.1
 Identities = 30/110 (27%), Positives = 40/110 (36%), Gaps = 23/110 (20%)
Query: 131 NEDTTSNTDETSNQNATSLDNS-----TGMTANRNAYVSLPQS----EVNIDVDNTTLRF 181
N +TT N ++N S DN+ T T N N + + D NT ++
Sbjct: 251 NINITQNLTTSTNTTTVSTDNNNNNINTKPTDNNNTNIKSTDNYNTGTKETDNKNTDIKA 310
Query: 182 ADMNTI------DNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                              DN KT S + SN + N K N
           DNN I
Sbjct: 311 TDNNNITITTDNTNTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVIST 360
>gi|1429240|emb|CAA67659| (X99260) lower collar protein
           [Bacteriophage B103]
           Length = 293
 Score = 43.8 bits (101), Expect = 0.001
 Identities = 53/204 (25%), Positives = 79/204 (37%), Gaps = 42/204 (20%)
Query: 56 EKVFKG----FSLKDELSDLLFKKSFTIHFLD----REINRQTVEAFGMQVITVCITHED 107
EK+ KG F + + D ++K F HF+ REI +T F + T I +
Sbjct: 26 EKIEKGRPKLFDFQYPIFDESYRKVFETHFIRNFYMREIGFETEGLFKFNLETWLIINMP 85
Query: 108 YLNVVYSSSEVEKY------LQSQGFTEH-----NEDTT------SNTDETSNQNA 146 Y N ++ S E+ KY L + G ++ N DTT SNT + NA
Sbjct: 86 YFNKLFES-ELIKYDPLENTRLNTTGNKKNDTERNDNRDTTGSMKADGKSNTKTSDKTNA 144
Query: 147 TSLDNSTGMTA------NRNAYVSLPQSEVNIDVDN--TTLRFADNNTIDNGKTVNKS 196
                            NR PS+N+ ++ TL+A + I+ T NK
                GT
Sbjct: 145 TGSSKEDGKTTGSVTDDNFNRKIDSDQPDSRLNLTTNDGQGTLEYA--SAIEENNTNNKR 202
Query: 197 SNESNQNAKRNQNQKGNAKGTQFT 220
                                                                              - _ ______
Sbjct: 203 NTTGTNNVTSSAESESTGSGTSDT 226
Query= pt|110879 44AHJDORF009 Phage 44AHJD ORF |5744-6496|2 1
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>gi|2764981|emb|CAA69021.1| (Y07739) N-acetylmuramoyl-L-alanine
           amidase [Staphylococcus phage Twort]
           Length = 467
 Score = 180 bits (452), Expect = 1e-44
 Identities = 89/157 (56%), Positives = 109/157 (68%), Gaps = 8/157 (5%)
         MKSQQQAKEWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFK 60
           MK+ +QA+ +I G DFDG YG+QCMDL+V Y+Y++TDGK+RMWGNAKDAINN F
         MKTLKQAESYIKSKVNTGTDFDGLYGYQCMDLAVDYIYHVTDGKIRMWGNAKDAINNSFG 60
Sbict: 1
Query: 61 GLATVYKNTPSFKPQLGDVAVYTNGQ---YGHIQCVLS----GNLDYYTCLEQNWLGGGF 113
G ATVYKN P+P+P+ GDV V+T G YGHI V + G+L Y T LEQNW G G
Sbjct: 61 GTATVYKNYPAFRPKYGDVVVWTTGNFATYGHIAIVTNPDPYGDLQYVTVLEQNWNGNGI 120
Ouery: 114 DGWEKATIRTHYYDGVTHFIRPKFSGSNS-KALETSK 149
              E ATIRTH Y G+THFIRP F+ +S K +T K
Sbjct: 121 YKTELATIRTHDYTGITHFIRPNFATESSVKKKDTKK 157
 Score = 61.7 bits (147), Expect = 6e-09
 Identities = 41/125 (32%), Positives = 57/125 (44%), Gaps = 8/125 (6%)
Query: 125 YYDGVTHFIRPKFSGSNSKALETSKVNTFGKWKRNQYGTYYRNENGTFTC-GFLPIFARV 183
           YY+G T
                           +K
                                  + +T G W N YGTYY++E+ TF C
Sbjct: 346 YYEGKTPV--PTVVNQKAKTKPVKQSSTSG-WNVNNYGTYYKSESATFKCTARQGIVTRY 402
Query: 184 GSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNWQGTR-YYLPVRQWNGKTGNSYSV 242
                             Y+ VC DGYVWI + G + ++PVR W+ N+ +
Sbjct: 403 TGPFTTCPQAGVLYYGQSVTYDTVCKQDGYVWISWTTNGGQDVWMPVRTWD---KNTDIM 459
Query: 243 GIPWG 247
Sbjct: 460 GQLWG 464
>gi|113675|sp|P24556|ALYS_STAAU AUTOLYSIN
           (N-ACETYLMURAMOYL-L-ALANINE AMIDASE)
           >gi|79887|pir||JQ1147 N-acetylmuramoyl-L-alanine amidase
           (EC 3.5.1.28) - Staphylococcus aureus >gi|153067
           (M76714) peptidoglycan hydrolase [Staphylococcus aureus]
           Length = 481
 Score = 118 bits (292), Expect = 6e-26
 Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)
Query: 135 PKFSGSNSKALETSKVNTFGK-WKRNQYGTYYRNENGTFTCGFLPIFARVGSPKLSEPNG 193
          P + SN + ++ V
                               WKRN+YGTYY E+ FT G PI R
Sbjct: 365 PVATVSNESSASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPITVRKVGPFLSCPVG 424
Query: 194 YWFQPNGYTPYNEVCLSDGYVWIGYNWQGTRYYLPVRQWNGKTGNSYSVGIPWGVFS 250
          Y FQP GY Y EV L DG+VW+GY W+G RYYLP+R WNG
                                                      + +G WG S
Sbjct: 425 YQFQPGGYCDYTEVMLQDGHVWVGYTWEGQRYYLPIRTWNGSAPPNQILGDLWGEIS 481
 Score = 78.0 bits (189), Expect = 7e-14
 Identities = 48/109 (44%), Positives = 62/109 (56%), Gaps = 6/109 (5%)
Query: 15 EGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDA-INNDFKGLATVYKNTPSFK 73
              + D YGFQC D + A + + G + AKD
                                                    N+F GLATVY+NTP F
Sbjct: 18 EGKOFNVDLWYGFOCFDYANAG-WKVLFGLLLKGLGAKDIPFANNFDGLATVYONTPDFL 76
Query: 74 PQLGDVAVYTNGQ---YGHIQCVLSGNLDYYTCLEQNWLGGGF-DGWEK 118
                       YGH+ V+ LDY EQNWLGGG+ DG E+
           O GD+ V+ +
Sbjct: 77 AQPGDMVVFGSNYGAGYGHVAWVIEATLDYIIVYEQNWLGGGWTDGIEO 125
>gi|1763243 (U72397) amidase [bacteriophage 80 alpha]
          Length = 481
                                                                             - _____
Score = 118 bits (292), Expect = 6e-26
Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)
Query: 135 PKFSGSNSKALETSKVNTFGK-WKRNQYGTYYRNENGTFTCGFLPIFARVGSPKLSEPNG 193
          P + SN + ++ V
                               WKRN+YGTYY E+ FT G PI R P LS P G
Sbjct: 365 PVATVSNESSASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPITVRKVGPFLSCPVG 424
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Query: 194 YWFQPNGYTPYNEVCLSDGYVWIGYNWQGTRYYLPVRQWNGKTGNSYSVGIPWGVFS 250
           Y FQP GY Y EV L DG+VW+GY W+G RYYLP+R WNG
                                                      + +G WG S
 Sbjct: 425 YQFQPGGYCDYTEVMLQDGHVWVGYTWEGQRYYLPIRTWNGSAPPNQILGDLWGEIS 481
 Score = 83.5 bits (203), Expect = 2e-15
 Identities = 50/115 (43%), Positives = 65/115 (56%), Gaps = 6/115 (5%)
           EWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDA-INNDFKGLATVYK 67
           EW+ EG + D YGFQC D + A + + G +
                                                   AKD N+F GLATVY+
Sbjct: 12 EWLKTSEGKOFNVDLWYGFOCFDYANAG-WKVLFGLLLKGLGAKDIPFANNFDGLATVYQ 70
Query: 68 NTPSFKPQLGDVAVYTNGQ---YGHIQCVLSGNLDYYTCLEQNWLGGGF-DGWEK 118
           NTP F Q GD+ V+ + YGH+ V+ LDY
                                                  EQNWLGGG+ DG E+
Sbjct: 71 NTPDFLAQPGDMVVFGSNYGAGYGHVAWVIEATLDYIIVYEQNWLGGGWTDGIEQ 125
>gi|4574237|gb|AAD23962.1|AF106851_1 (AF106851) LytN (Staphylococcus
           aureusl
           Length = 383
 Score = 84.3 bits (205), Expect = 9e-16
 Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)
Query: 15 EGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
           E G DFDG+YG+QC DL Y ++ +++G
                                                    N+F A +Y NTP+FK
Sbjct: 252 ENRGWDFDGSYGWQCFDLVNVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311
Query: 75 QLGDVAVYT---NGQYGHIQCVLSGNLD----YYTCLEQNWLGGGFDGWEKATIRTHYYD 127
+ GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A H Y+
Sbjct: 312 EPGDLVVFSGRFGGGYGHTAIVLNGDYDGKLMKFQSLDONWNNGGWRKAEVAHKVVHNYE 371
Query: 128 GVTHFIRP 135
Sbjct: 372 NDMIFIRP 379
>gi|3767593|dbj|BAA33856.1| (AB015195) LytN [Staphylococcus aureus]
           Length = 383
 Score = 84.3 bits (205), Expect = 9e-16
 Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)
Query: 15 EGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
           E G DFDG+YG+QC DL Y ++ ++ +G
                                                    N+F A +V NTP+PK
Sbjet: 252 ENRGWDFDGSYGWQCFDLVNVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311
Query: 75 QLGDVAVYT---NGQYGHIQCVLSGNLD----YYTCLEQNWLGGGFDGWEKATIRTHYYD 127
           + GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A
Sbjct: 312 EPGDLVVFSGRFGGGYGHTAIVLNGDYDGKLMKFQSLDQNWNNGGWRKAEVAHKVVHNYE 371
Query: 128 GVTHFIRP 135
               FIRP
Sbjct: 372 NDMIFIRP 379
>gi|2764983|emb|CAA69022.1| (Y07740) cell wall hydrolase Ply187
           [Staphylococcus phage 187]
           Length = 628
 Score = 76.9 bits (186), Expect = 2e-13
 Identities = 50/144 (34%), Positives = 68/144 (46%), Gaps = 18/144 (12%)
          QQAKEWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMW-----GNAKDAINNDF 59
           +Q +W G+GVD DG YG QC DL Y++ R W GNA+D
Sbjct: 12 KQVVDWAINLIGSGVDVDGYYGRQCWDLP-NYIFN-----RYWNFKTPGNARDMAWYRY 64
Query: 60 KGLATVYKNTPSFKPQLGDVAVYTNGQY----GHIQCVLS-GNLDYYTCLEQNWLGGGF 113 V++NT F P+ GD+AV+T G Y GH V+ Y+ \frac{1}{2} ++QNW
                                                                                  ____
Sbjct: 65 PEGFKVFRNTSDFVPKPGDIAVWTGGNYNWNTWGHTGIVVGPSTKSYFYSVDQNWNNSNS 124
Query: 114 DGWEKATIRTHYYDGVTHFIRPKF 137
                    H Y GVTHF+RP +
               A
Sbjct: 125 YVGSPAAKIKHSYFGVTHFVRPAY 148
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>gi|3287732|sp|005156|ALE1_STACP GLYCYL-GLYCINE ENDOPEPTIDASE ALE-1
           PRECURSOR >gi|1890068|dbj|BAA13069| (D86328) ALB-1
           [Staphylococcus capitis]
          Length = 362
 Score = 73.4 bits (177), Expect = 2e-12
 Identities = 47/117 (40%), Positives = 61/117 (51%), Gaps = 10/117 (8%)
Query: 132 FIRPKFSGSNSKALETSKVNTFGKWKRNQYGTYYRNENGTFTCGFLPIFARVGSPKLSEP 191
                GSNS TS N G +K N+YGT Y++E+ +FT
                                                      T R+ P S P
          F++
Sbjct: 252 FLKSAGYGSNS----TSSSNNNG-YKTNKYGTLYKSESASFTAN-TDIITRLTGPFRSMP 305
Query: 192 NGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYYLPVRQWNGKTGNSYSVGIPWG 247
                     Y+EV DG+VW+GYN G R YLPVR WN TG +G WG
Sbjct: 306 QSGVLRKGLTIKYDEVMKQDGHVWVGYNTNSGKRVYLPVRTWNESTG---ELGPLWG 359
>gi|79926|pir||A25881 lysostaphin precursor - Staphylococcus
          simulans >gi | 153047 (M15686) lysostaphin (ttg start
          codon) [Staphylococcus simulans]
          Length = 389
 Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)
WK N+YGT Y++E+ +FT
Sbjct: 258 HFQRMVNSFSNSTAQDPMPFLKSAGYGKAGGTVTPTPNTGWKTNKYGTLYKSESASFTPN 317
Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYYLPVRQWNG 234
I R P S P + Y+EV DG+VW+GY G R YLPVR WN Sbjct: 318 -TDIITRTTGPFRSMPQSGVLKAGQTIHYDEVMKQDGHVWVGYTGNSGQRIYLPVRTWNK 376
Query: 235 KTGNSYSVGIPWG 247
           T ++G+ WG
Sbjct: 377 STN---TLGVLWG 386
>gi|126496|sp|P10548|LSTP_STAST LYSOSTAPHIN PRECURSOR
           (GLYCYL-GLYCINE ENDOPEPTIDASE) >gi|79927|pir||S01079
          lysostaphin precursor - Staphylococcus simulans bv.
          staphylolyticus >gi|581744|emb|CAA29494| (X06121)
          lysostaphin (AA 1-480) (Staphylococcus simulans bv.
          staphylolyticus)
          Length = 480
 Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)
Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK------WKRNQYGTYYRNENGTFTCG 175
          HFR S SNS A + K +GK
                                                WK N+YGT Y++E+ +FT
Sbjct: 349 HFQRMVNSFSNSTAQDPMPFLKSAGYGKAGGTVTPTPNTGWKTNKYGTLYKSESASFTPN 408
Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYYLPVRQWNG 234
            I R P S P
                            + Y+EV DG+VW+GY G R YLPVR WN
Sbjct: 409 -TDIITRTTGPFRSMPQSGVLKAGQTIHYDEVMKQDGHVWVGYTGNSGQRIYLPVRTWNK 467
Query: 235 KTGNSYSVGIPWG 247
           т
               ++G+ WG
Sbjct: 468 STN---TLGVLWG 477
>gi|3287967|sp|P10547|LSTP_STASI LYSOSTAPHIN PRECURSOR
          (GLYCYL-GLYCINE ENDOPEPTIDASE) >gi 2072411 (U66883)
          lysostaphin (Staphylococcus simulans)
          Length = 493
Score = 69.5 bits (167), Expect = 3e-11
Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)
Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK------WKRNQYGTYYRNENGTFTCG 175
          HPR SSNSA+ K +GK
                                                WK N+YGT Y++E+ +FT
Sbjct: 362 HFQRMVNSFSNSTAQDPMPFLKSAGYGKAGGTVTPTPNTGWKTNKYGTLYKSESASFTPN 421
Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYYLPVRQWNG 234
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DG+VW+GY
                                      Y+EV
              I R
                    PSP
                                                        G R YLPVR WN
Sbjct: 422 -TDIITRTTGPFRSMPQSGVLKAGQTIHYDEVMKQDGHVWVGYTGNSGQRIYLPVRTWNK 480
Query: 235 KTGNSYSVGIPWG 247
                ++G+ WG
Sbjct: 481 STN---TLGVLWG 490
>gi|3341932|dbj|BAA31898.1| (AB009866) amidase (peptidoglycan
          hydrolase) [bacteriophage phi PVL]
          Length = 484
 Score = 68.3 bits (164), Expect = 6e-11
 Identities = 52/150 (34%), Positives = 71/150 (46%), Gaps = 17/150 (11%)
          SQQQAKEWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGL 62
                     G + D YGFQC D + + + I G+ R+ G
                                                            IDK
          TKNQAEKWFDNSLGKQFNPDLFYGFQCYDYASMF-FMIATGE-RLQGLYAYNIPFDNKAR 61
Sbjct: 4
Query: 63 ATVY----KNTPSFKPQLGDVAVYTN---GQYGHIQCVLSGNLDYYTCLEQNWLGGGF-- 113
Y KN SF PQ D+ V+ + G GH++ V S NL+ +T QNW G G+
Sbjct: 62 IEKYGQIIKNYDSFLPQKLDIVVFPSKYGGGAGHVEIVESANLNTFTSFQNWNGKGWTN 121
Query: 114 ---- DGW-- EKATIRTHYYDGVTHFIRPKF 137
               GW E T HYYD +FIR F
Sbjct: 122 GVAQPGWGPETVTRHVHYYDDPMYFIRLNF 151
Query= pt|110882 44AHJDORF012 Phage 44AHJD ORF |8391-8813|3 1
         (140 letters)
>qi|140528|sp|P24811|YOXH BACSU HYPOTHETICAL 15.7 KD PROTEIN IN
          SPOILIC-CWLA INTERGENIC REGION (ORF2)
          >gi|322189|pir||B44816 orf2 5'of autolytic amidase -
          Bacillus subtilis >gi|142801 (M59232) open reading frame
          2 [Bacillus subtilis] >gi|1217874|dbj|BAA06959| (D32216)
          ORF121 [Bacillus subtilis] >gi|1303767|dbj|BAA12423|
          (D84432) YqdD [Bacillus subtilis]
          >gi|2635036|emb|CAB14532| (Z99117) alternate gene name:
          yqdD; similar to holin (Bacillus subtilis)
          Length = 140
 Score = 80.4 bits (195), Expect = 6e-15
 Identities = 45/130 (34%), Positives = 67/130 (50%), Gaps = 3/130 (2%)
          VKFRFTDSEAFHMFIYAGDLKLLYFLFVLMFVDIITGISKAIKNNNLWSKKSMRGFSKKX 63
Query: 4
          + F D ++F G +K L L VL +D++TG+ KA K L S+ + G+ +K
          INFETLDLARVYLF---GGVKYLDLLLVLSIIDVLTGVIKAWKFKKLRSRSAWFGYVRKL 64
Sbict: 8
G L T+ +YIANEGLSI EN A++ V +P I D+L+ I
Sbjct: 65 LNFFAVILANVIDTVLNLNGVLTFGTVLFYIANEGLSITENLAQIGVKIPSSITDRLQTI 124
Query: 124 KNDTEKSDNN 133
          +N+ E+S NN
Sbjct: 125 ENEKEQSKNN 134
>gi|4126631|dbj|BAA36651.1| (AB016282) ORF45 [bacteriophage phi-105]
Score = 76.1 bits (184), Expect = 1e-13
Identities = 44/115 (38%), Positives = 61/115 (52%), Gaps = 4/115 (3%)
Query: 21 GDLKLLYFLFVLMFVDIITGISKAIKNNNLWSKKSMRGFSKKXXXXXXXXXXXXXXXXXXXXXXXXXXXX
          G++K L + VL +DIITG+ KA K L S+ + G+ +K
Sbjct: 17 GEVKYLDLMLVLNIIDIITGVIKAWKFKELRSRSAWFGYVRKMLSFLVVIVANAIDTIMD 76
Query: 81 XKGGLLMITIFYYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKND----TEKSD 131
                                                                           - _
            G L T+ +YIANEGLSI EN A++ V +P I D+L VI++D
                                                         TEK D
Sbjct: 77 LNGVLTFATVLFYIANEGLSITENLAQIGVKIPAVITDRLHVIESDNDQKTEKDD 131
>gi|141088|sp|P26835|YNGD CLOPE HYPOTHETICAL 14.9 KD PROTEIN IN NAGH
          3'REGION (ORFD) >gi|1075967|pir||S43905 hypothetical
          protein D - Clostridium perfringens >gi 455154 (M81878)
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ORF D (Clostridium perfringens) Length = 132

Score = 60.9 bits (145), Expect = 4e-09 Identities = 38/127 (29%), Positives = 63/127 (48%), Gaps = 3/127 (2%)

MNEVKFRFTDSEAFHMFIY-AGDLKLLYFLFVLMFVDIITGISKAIKNNNLWSKKSMRGF 59 +I+ A D+ L+ L V +F+D +TG+ K K+ L S +N +K+

INYIKWGIVSLGTLFTWIFGAWDIPLITLL-VFIFLDYLTGVIKGCKSKELCSNIGLRGI 63 Sbjct: 5

I ++YI NEG+SI+ENCA + V +PE++K Sbjct: 64 TKKGLILVVLLVAVMLDRLLDNGTWMFRTLIAYFYIMNEGISILENCAALGVPIPEKLKQ 123

Query: 119 KLRVIKN 125 L+ + N

Sbjct: 124 ALKQLNN 130

>gi|2293160 (AF008220) YtkC [Bacillus subtilis] >gi|2635548|emb|CAB15042| (Z99119) similar to autolytic amidase [Bacillus subtilis]

Length = 134

Score = 36.4 bits (82), Expect = 0.099 Identities = 25/109 (22%), Positives = 41/109 (36%)

Query: 17 FIYAGDLKLLYFLFVLMFVDIITGISKAIKNNNLWSKKSMRGFSKKXXXXXXXXXXXXXX 76

L LM ++ I+ K + L KK F + G KK

Sbjct: 20 FFFGGFQYSFLILLSLMAIEFISTTLKETIIHKLSFKKVFARLVKKLVTLALISVCHFFD 79

Query: 77 XXXXXKGGLLMITIFYYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKN 125 +G + + I +YI E + IV + + + VP+ + D L +KN

Sbjct: 80 QLLNTQGSIRDLAIMFYILYESVQIVVTASSLGIPVPQMLVDLLETLKN 128

>gi|1181973|emb|CAA87743.1| (Z47794) holin protein [Bacteriophage

Length = 134

Score = 31.3 bits (69), Expect = 3.3 Identities = 27/88 (30%), Positives = 36/88 (40%), Gaps = 5/88 (5%)

Query: 29 LFVLMFVDIITGISKAIKNNNLWSKKSMRGFSKKXXXXXXXXXXXXXXXXXXXXXXXX 86 G +L

LF L+ D ITG KAK S ++G K

Sbjct: 18 LFALILFDFITGFLKAWKWKVTDSWTGLKGVIKHTLTFIFYYFVAVFLTYIHAMAVGQIL 77

Query: 87 MITIFYYIANEGLSIVENCAEMDVLVPE 114

++ I YA LSI+EN AMV +P+
Sbjct: 78 LVIINLYYA---LSIMENLAVMGVFIPK 102

Table 21

# Phage 182 complete genome sequence. 17833 nucleotides.

1	tagaatattg	tcataaaaca	casacataat	aatocatatt	attotttaca	aatatotaat	ttcgtgatat	
	-agaacacty	taagttaaag	caaacacaac	22022022	cateaatact	tterseattr	cassactat	
71								
141	tggaggaaaa	ataatgaaat	atteactaca	acadalayat	gaaattaaat	caacaacccc	cagaaccaga	
211	ttaaaaaggc	atgaactaga	ggaattggtg	gacgaagtaa	acgatattgc	taaagateeg	gaggaaagat	
281		gttttattac						
351		aagatcacaa						
421	aaataattac	acaaaaagct	ttacaaatat	aacacatcat	gttatactaa	aagagtagta	agggaacgga	
491	aaatacctta	cttcacacct	caatcattct	tatcaaaata	caaaaggagg	gaaaataatg	ggtcgaaaac	
561	taatqcaacq	aaacgtaaca	tcaactaaag	tagaattctc	agaagttatc	gtacaagatg	gagegeeaac	
631		tgcgaaccag						
701		ctgataaaaa						
771		tatcgagtta						
841		atcatggaaa						
		caaacggggc						
911								
981		ctcagacgaa						
1051	_	gatggtaaaa				_		
1121		ctgctaaccc						
1191		aaatctacaa						
1261		ggtggttttt						
. 1331	aatgatagag	ccaagttaga	gaaaatctac	ggtaaatcta	acaaagctcg	taaaaaatac	aatcgtttaa	
1401	gacaaaaagg	agttgaggaa	aggcaacttc	caactgttcc	aacatcaaag	aaaagactta	ttgactacgt	
1471	aaaatcaaca	aatatgagtc	gtagtgattt	taacaagatg	ttagacgagt	tggtagattt	tgcacaacct	
1541		attacatttt						
1611		aacagagcaa			_			
1681		acagaaaaca						
1751		cagattttaa						
1821		agacgaacaa						
1891		ttcaattcag						
1961		atgttagtaa						
2031		acaagtttac						
2101		acgaagaaca						
2171		aacaacaact						
2241		atgacgttcg						
2311		tccacaacga						
2381		agaagcaaaa						
2451		tgttgggaag						
2521		atgatagcct						
2591		aggcgaaata						
2661		aagaacgaca						
2731	cgaatgacta	gaggaagcga	cgctttaggc	gattacaaag	attggctaaa	agctacacat	ggaaaatcaa	
2801	ctttcaaaca	atggtttcct	attttgtctt	tagggtttga	taaagactta	cgtaaagcat	acaaaggcgg	
2871	cttcacttgg	gtaaacaaag	tttttcaagg	gaaagaaata	ggtgacggca	ttgtctttga	tgtcaactct	
2941		ctcaaatgta						
3011		cgactatccg						
3081		caagttaagc						
3151		acgaattaat						
3221		gatacattac						
3291		atcgaagtaa						
3361		gaaagttcgg						
3431								
3501		gacactagga						
		agatatacta						
3571		atctagtagg						
3641		gcatgaaagc						
3711		ttaaatgtaa						
3781		gtttttcaag						
3851		tacaatcaaa						
3921		attgacggtt						
3991	tatggaaaag	aaacacgtga	aattgaagca	gtaacattag	taaaaacagg	aaatttaaaa	aaataaatta	
4061		ttgcaaagta						
4131	ggcgaatgta	cacgtgaaat	atcgtgcgct	cccgttaagt	tatggacaca	taaacgtttt	gaccgtcaac	
4201		aaccttttag						
4271	_	acagttttta						. –
4341		taaatggtat						
4411		gcgagaagac						
4481		gataacgaag						
4551	2ccccaaaa	ataacgaacc	agaaacagac	cagaatatta	cactagacga	tttaggaatt	taaggaggaa	
4621	2222025000	tgacaaaatc	2200000000	atottettee	toccacaaat	ataggaacac	cagtacaatt	
4621	aaaatatyyC	atttataata	attestests	tettttes	acasacatea	ctatoccasa	tacageteec	
	GALMACLUCE	actialaded	guerate	LLLLLLLAY	2-22		-3005acaac	

atcgaagcgg ttggtgcagg gatcacacgt ttagacgtag taaaaaacga atttatttca actttagttg accgtattgg taaagtagtt atccgataca aatcttggcg taaccctttg aaaatgttta aaaaaggaaa 4761 4831 catgccttta ggtcgaacga ttgaagaaat ttttgttgac attgcacagg aacataagtt caaccctgac 4901 gagtotgtta caggggtatt taaacaggaa gttoccgatg taaaaacatt gttocacgaa attaatcgtg 4971 aaggttacta caaacaaacg atccaagaag catggttaga aaaagcattt acttcatggg ataatttcaa 5041 tagtttegtt getggtgtaa tgaaegettt atacacaggt gaegaagtaa gegaatttga atacaegaaa 5111 ttattaatag caaactacca agaaaaagag ctattcaaag agatcgaaat tggcgaaatt actgaatcaa 5181 atgcaaaaga atttatccgt aagatcaaat caacctctaa caaattagaa tttatgagtt ccgcttacaa 5251 cgctcaagga gttaaaacat ctacctcaaa atctgatcaa tacgttatta ttgacgccga cacagacgca 5321 accattgacg ttgacgtttt agcagcggca ttcaatatga gtaaaactga ctttgtagga cacaaaatcg 5391 5461 ttattgatga gtttcctaaa aaagaaggcg aagaatcgtc aaatattgtg gcagttattg tagatagtga atggtttatg atctacgaca aattgtacaa aacaacaagt ctatacaacc ctgaagggtt atattggaat 5531 tattggttgc accaccacca actatattct acttctcaat tcgggaacgc tgttgctttt gttaaatcag 5601 caacaaaacc tgtcacaaaa gttgcttttg caagtgcaac aactagtgtt gttaaaggat catctaaaga 5671 tatcgcattg acatttacac cagtagaagc aacaaaccaa caaggagaag ttgtttcatc agcaccagca 5741 ttggttaagg caaccgtaaa acaaacagca ggtaaagcga ctgccgtaac cgtagaaggc ttagaagtcg 5811 5881 gtcaatcatt agtaacattc acagetatcg gaggtcaaca agcaacggtt cttgttacgg ttacttctga ctaaggagga caattatggc aagaaggtat acaaatgtaa aattgttggc taacgtgcct tttgataaca 5951 cctatacaca cacaagatgg tttaaaactc aacaggaaca ggaatcgtac tttaattcgt ttcctgttct 6021 taacgagaat agagattgtt cttatcaaag ggatacacaa ctcgggggag tttttagagt agataaacac aaagacgcct tatatgcttg taactatctc atctttaaaa acgaagaaac ttatcctagt aaatggcagt 6091 6161 atgeetttgt tactgatatt gaatataaga atgacaacac aagtttegtt acetttgaaa ttgatgtttt 6231 acaaacttat cgtttcgata ttggtatacg agaaagtttc attgcaaaag aacaccctca actttattat 6301 toqaatqqaa tacctttcat taatacaatt gaagagtogo ttgattacgg tagagaatac acaacaacaa 6371 atqtaacaac ttttcatcct aacgatggag tcaattttct tgttattcta acaagtgaag caatgccagt 6441 tggagataag gaagataaat caggaggatc aatagtaggt ggcccatctc ctttttccta ttatttactt 6511 cctatcaatt caagtgggga ggtatacaaa ccaaatgggg caggcaatgc taattttgga gagtacatgg 6581 cgtttcttac aacgaaagaa ccttttttaa ataagatagt cgggatgtat gtaacgtcgt atacaggtat 6651 accattcatt gtggatcacg cgaacaaaac ggtaaggtat aatgcaggag gttcttataa gatcatgctt 6721 ccaacctacg ctagtgatcc aacaggaaca atgaaaacat tcgctttctt ttgtgtaaaa gaagcaagaa 6791 6861 cattegtace taaaagaatt gatettgtag ggaacgtgta taactacttt agagaagett tteegtttaa tottaaqqaa tcaaaactat ttatgtatcc ctattgttta atagaaatta cagatacaaa aggacatgta 6931 atgactttaa gacctgaata tottacaggt ggtaaattga gtgtatatgt aaaaggttcg ttaggaattt 7001 ctaataaagt gatgatcgag ccgattgatt atgatgtaag taactcaacc attattacca atttaagtga 7071 caagatgtta atcgataatg atcctaacga tgtaggagtt aaatctgact atgcttctgc attcatgcaa 7141 ggaaacaaaa actccttgat tgctcaagag caaaacattc gcaatacttt cagacatggt atgggaaaca 7211 gtgcaatgag tacaggagga gcgatctttt cagccttagc aagtaacaac ccttttgttg gtttgactaa 7281 catcatggga gcaggacaac aagtaaacaa ctatgtttct gaaaaagaaa acggtttgaa cctcttggca 7351 ggtaaagtgg cagatatcga aaatattcca gataatgtaa cacagettgg atcaaactta tetttcacaa 7421 caggaaactt tcaaaactat tatcaattgc gcttcaaaca aattaaatat gagtatgcaa caagacttga 7491 tegttactte teaatgtatg geacaaagag caategagta getacaccaa aettacaaac aagaaaagca 7561 tggaatttca ttaaattaaa agaaccaaat attgtaggca caatgagtaa cgatgtatta acacgtgtga 7631 aacaaatttt tagtgcaggc gttacgcttt ggcatacgaa tgatgttttg aattataacc aagacaacgg 7701 agatgtatag gaaggaggaa taagatgagt agacgaaaag gtgcaggact tgctagaaat aaccgttata 7771 7841 cagcaaaaag cagacettat ccaaatgaac cctattcaag tgatgtagaa gaaatcaget actatgaaca ttatcgtaga caactcacgc tccttacgtt tcagttgttt gaatgggaaa atttgccaaa atcaattgac 7911 cotegitatt tagaaattge tttacacact aatggttate ttggtttett taaagaccet acacttgggt 7981 tcatggtttg cgcaggggca gaagatggtc aaatcgatca ttatcacaac cctattttct ttacagcaaa 8051 cgaagcaatg tatcacaaga gatatcctgt tttaagatat gatgatgatg atgataaatc aaaatgtatc 8121 atgttgtata ataatgactt gaaagtteet acgttaccaa gtttacatcg ttttgcttta gatatggcgg 8191 acataaacca gatatcacga gtgaatcgaa gagcgcaaaa aacacctgta attattcaaa ctgatgaaaa 8261 gaaatacttc tcattgctac aagcttataa ccaaattgac gaaaataatc aggctgtttt tgtggataaa 8331 gatatggagt ttgacgaatc ttttaatgta tggcaaacaa atgctccata tgtagtagat aaactacgat 8401 cagaattgaa cgaagtatgg aatgaagtgt taacttttct aggtatcaac aatgctaacg tagataagac 8471 tgcacgtgta caaacatcag aagtettate taacaatgaa cagattgaaa gttcaggtaa catettgtta 8541 aaatcaagaa aagagttttg cgatcgtgta aatcgtgtct ttggcgatga acttgacgga aagattgacg 8611 tgaagtttag aacagacgcc gttcgacaat tacaactggc ggcaggtcaa tcaaaaaaag accagatgag 8681 tggagggttg ccaagtgcta cttaaacgtt atattgaaag tttcacttat taccaacctg aattatctcg 8751 8821 aaaagaacgt attgaagttg gccgaaaaca attgtttgat tttgattatc cgttttatga cgaaacaaaa cgagcagaat ttgaaacaaa atttatcaat cacttttact tgagagagat aggctcagaa acgatgggat 8891 catttaagtt taatettgae gaatatttaa atetaaacat geeetattgg 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tggtacgtta gaaaatttaa tcaatgatac tgtttttgca aattatatca aagaaatcaa aagattacaa atcttggttg 9871 ctgaaacacg tgctaacagt gtgaatattc ttttgacaaa aaataaaccg gatgttgctg atgatcgaac 9941 attttggtat aagattcaac gcgacaatac tgattatgga gccgatccta ttgacacgtt acgtattgtt 10011

10081 gcaatcaata aagttagtgg ctggaatacc gctacaggag atatttatct taacattaaa ggaacggagg 10151 gtgtataatg gcagacatta gaacacaact aacaagtgaa gatggatcag acaatttatt tccaatttca 10221 aaagccgtta atattatgac taatagcggt acgaatgtag aaggagaatt gggtacactc aaacaaaatg 10291 acgaaacaat gaatacctca gttcaaaatg ctgtagttac tgccaatcaa gcaaaagatt ctgtagctga 10361 attaaatgta aatgttggta aactaaccaa tcgaataaca acattagaga gtacagtggc taatcttgat ggtattcgtt atgtagaggt gtaatatggc agataaaaat attcaaatgc aggataaaga tcataatcgt 10431 10501 ttaatgcctg ttacaattgc taaaaatgtt ctaacaggcg actctaatct tgaattagtt aatgctgaaa 10571 taagaggtaa cgctagtgaa gctaaaacac ttgcacaaca agctaaagaa actgctgctg gtttgtcaac 10641 agaaattgac acagtaacat caaccgcaaa tcaagcgttg acgaaggctg gtacagcaca acaaaccgca 10711 gaacaagega aaacaacage aaacagtate agegeagttg caacggeage taaaaacaca getgatteag 10781 cacaaaaaag tgcaactgat ctagctgttc gagtaagcag tttagaggac acagcaatac aatatactgt attaccatag gaggaaaaat aatggcaaat aaaaatattc aaatgaagga tagcaatgac aataatttat 10851 atccaagtgt tcgagcagaa aacttgttag atttgaccag tcgtgctgaa ttaacaatga caaattgtca 10921 10991 attatatgca gctggtgata aaacaaatgc aatctcttat ctcggtgcag taggtatgct cgaaggtatg 11061 ataaagttta ctgaaagttt gacaaaccct gtgatcacaa cgctaccaga aggttttaga ccaataagaa caaaacgtat tggttgtttc gcaaaatatt acacaccaaa tccaacagat acaaaagaaa tggtttatgt 11131 atcaatcaca cctgatggca aagtaactgt aaatgacaat gtaggtaaaa tcgaatatct atccctagat 11201 11271 aattgegttt teeetetaaa ataaggaggt teatatggaa gaacgaattg atatteaaat gaacaagatg aaagaagaaa atcaaaagaa ttacctattg caccctgaaa cgaacccgaa acaagttgtt tttgatgaaa 11341 cattgcatgg aaatgaaaat caggagagtt tcaacaattt tgttgacaca agaaaaatga caactacaat 11411 11481 tgatgtaagt gcttatgggg ttatcgctga cggtgtaaca gattgtacac caatattaaa taaattactt gaagaaaaaa gcgaaatggg tatcactttt tattttcctc cttgtgaacg tgattcatat tatcgctttg 11551 11621 ctaacaccat tgaattgaaa cgtgatgtac ctgtagttac tttcttagga tcgggagaaa cgacattaaa 11691 gtttgaaaca atgacggcat ttaatgtaaa catcgaaagt ttcaatattg atggttttgc attatggttg ccacaaggcg ctcaaagtgg taaaggaatt ttctttaatg atactegcaa ttacaatcgt tttgactttg 11761 11831 atttgtttgt tcgtaactgt actttaaatg aaggaacgta tgttgttgtt gctagaggta gaggggttac 11901 atttgaaaat tgtctattct ctaatatctc tcaagcaatt atcaaaacag cttttcccga tgtaaatggt 11971 atgtggcaag ggaacgatat caatactagg ggtacaggtt ttagaggttt ctttgtgaaa aacaaccgta 12041 ttcatttttg tacagegate attategaca atgacgatga ttateagaat gtaattaatt tetgtgaaat ttctggtaac acaatcgaag gtggcgtaag ttattatcga ggatatgcgc ataacttgca tgtccaaaac 12111 aacaaccatt ttctagcata cggaaataga aacgctttgt ttgagtttca agatgtggat caagcttata 12181 12251 ttgatgtaga tgtttattgt cgtaactcac aagtcgaggg aatgaatagt acagctattt cacgtttaat 12321 tgttgtttac ggacattacc gaaacttaaa gattacaggt aaattatatc gttgtcaagg acatgttatc acgttgtatg gcggtggcgt taatttctat tgtgacttga tggcacaaga agcacctttg acggacggtt 12391 12461 accegettat tcaaaceget gacaatcgag ttaactatga tgggtttgtt gttcgtggtt tgtctaattc 12531 aacaaaagta aatacaccaa tgatctataa agcacctcag actgttttct ataatcgtag aatcgatcat 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13441 ttggcgacaa agttaagaac ggacaagttt gcgcaatacg tgacgcggat catttacatt taggttttac 13511 taaaaaagat tttatgactg cgttaggatc ttctttcata gatgatggaa catgggaaga ccctttgaag 13581 tttttagggc aatgttttgg agatggagat actggcggag ataatgacga taacaataag gataaaaatg 13651 atettattta tetattgeta teegatgeet tgaatggttg gaaattttaa taaggagaaa aaggtatgat 13721 agaatatatc acacaatggt tggcagatga taatcatctt gtttatggtt tgattatatg gttaatggtt gcaatgatta tcgattttgt gttaggtttt acaattgcca aatttaacaa ggaaatcgac tttagtagtt ttaaagctaa agcaggtatc attgttaagg tggcagaaat ggtttagtg gtttacttta ttcctgtagc 13791 13861 13931 agtaaaattc ggtgcagtag gtattacaat gtatataaca atgttggttg gtttgatttt atcagaaatt 14001 tatagtatac taggacatat ttcagatatc gatgatgata ataattggac tgattatgtt aagaagtttt 14071 tagacggaac actcaacaga aaggacgata ttaaatgatg aatggtattg atatctctag ttatcaaaca 14141 ggaattgatc tttcaaaagt tccatgcgat tttgtaaata ttaaagcaac aggcggaaca ggttatgtaa 14211 accetgattg tgacegagea tttcaacaag ctttgtcttt aggtaaaaag attggtgtgt atcattttgc gcatgagagg ggtttagaag gtacacctca acaagaagcg caattetttt tagataatat taagggttacattggtaaag ctgttettat tettgacttt gaagggtcaa atcagaaaga tgtaaattgg gcgaaagcat 14281 14351 14421 ttcttgatta tgtttataat aaaacaggcg ttaaagcatg gttttatacg tatacagcaa acctcaatac 14491 aactgatttt tctagtattg caaaaggoga ttatggttta tgggttgctg aatatggatc aaatcaacca 14561 caaggetact etcaaccage gecacetaaa acaaataatt ttecaattgt tgeetgtttt cagtttacaa 14631 gtaaaggacg tttaccagga tacaacggca atcttgattt gaatgttttc tatggcgatg gtaatacatg 14701 ggatctgtat gtaggtaaaa aacaggatca aattgttcct cctgaaaata aaatatttga cgccacaagt 14771 gatgagttta ttttcactct tacaacaggt agcacaagcg tgttttattt tgacggagaa acgatctttg 14841 aattgtotga tocaacacaa otogatcata ttagaggaac atacaatcat gttcatggaa aagaaatcoc 14911 atcaatggtg tggacacctg aacaatttga tatttactta aaaatgtatg aaaagaaacc agtatataaa 14981 taggagtgta tagtatgaca aatagettag gegttaaaet tgaagagaaa aaettataet ataaceetaa 15051 caatgettta ggttttaatt geetaatgtt gtttgtaata ggegeaegtg gtataggtaa aacttatggt 15121 tataaaaaat ttgttgttaa tcgctttatt aaacacggcg aacaatttat ttatttaaga agattcaaaa 15191 cagaacttaa aaagatteet caatttttea aaacaatgge gaaagaattt eetgateata aacttgaagt 15261 aaaaggaaaa gaattctatt gtgatgataa attaatgggt tgggctgttc cacttagtac gtggggaatt gaaaaatcta atgaatatcc cgaagttcgt acaattttgt ttgatgagtt tttaattgag aaatcaaaaa 15331

15401	tcacttattt	accaaacgaa	gctgaagcct	tattgaacat	gatggaaacg	gttttccgaa	gacgtacaaa	
15471	tacaagatgt	gttatgttga	gtaatgcaac	tagtgtagtg	aacccttatt	tcttgtattt	caatctgcag	
15541	ccagatttga	ataagcgttt	taatctatat	caagatcgag	gtatattgat	tgaattgtgt	gattcaaaag	
15611	actttgcaga	agtgaagaga	gaaacacctt	ttggtagatt	gattcgtgga	acagaatacg	aagattttag	
15681	tatcaacaat	gagtttgtca	atgatagtga	tacgtttatt	gaaaagagaa	gtaaaaatag	tagtttctta	
15751	tgcgccattg	cttttgaagg	gaaaatcttt	gggtattgga	tagacgctga	aacaggttgt	gtctatgtga	
15821	gttatgatta	tcaaccaaat	acaaatcatt	tttatgcaat	gactacgaaa	gaccatgaag	aaaatagatt	
15891	gctgatgaaa	aattggcgaa	ataattatta	tctttcaaca	gtggcgaaag	cattcaagaa	tagttatctg	
15961	cggtttgata	acattgttat	taagaattta	cattatgatt	tgtttaataa	gatgaaaatc	tggtaaccct	
16031	attttagtag	agctaccacg	attagttcta	ttacaatgat	gaatagtaga	taacatagta	attgtagtct	
16101	gcgatagttt	tgttttggtt	ctttggcgtt	agtgattttt	gctaacgcct	ttttgtttgc	ttttggatcg	
16171	ggtgtgttaa	tgtagacgaa	atcttttctc	atagttcttt	ctccttatac	agttttaata	attccctgta	
16241	aaatgtagct	ataggacgtc	catttcttc	tattctaacg	caattcacta	tatccatttc	taggtatata	
16311						caaaacctcc		
16381	tataaaatac	tgtgatatcg	tatattggtt	ccttgtagaa	tgtagccatt	attccacctc	ctttaaatag	
16451	ccttttggta	tttgtaacgc	taactgatag	cgagaaccaa	cttttacgta	tgaagttact	aatttcattg	
16521	cctgacaata	cttttcaaga	atgttaaatt	gactcgattc	gggtaatagc	gttgaatgag	ttaacaaaag	
16591	ttcggtgata	tttatttccg	gaacgtcgaa	atcttgtaaa	gtcccctcta	tgatctctat	tttttcattg	
16661	tctgaaaggt	tacgtttaca	gtagaaacgt	aaccattcaa	ttagttcgcg	gtgttctttg	aatgttcgtg	
16731	caatcatttt	aattcctcct	atttgtccgt	aatttgttta	tatccgtcat	gtttcaattg	ttccgcatag	
16801	tgttcaacgc	ttttcattga	tttcgttatt	gcgatattaa	tgcaatggct	atcaagataa	acatagttat	
16871	atttatcatg	tgttaacacg	aactcttttg	taacgtaatc	aatgtataaa	attaattgtt	ttcctccttg	
16941	tgttatttct	gacttgatag	acgctaaact	atcgttgtca	tctttagtta	gttgatttaa	accctctaaa	
17011	attaatgata	aattgttaat	catgtaaaac	actcctttta	tattaatttg	atattgatac	caccaatcga	
17081						catgaatact		
17151						catcgcctac		
17221	tcaataagat	aatgtttatt	gttttcggta	tctatgatat	gataattcat	atcccactca	ttaaaggggt	
17291	gaagtagaga	tacctctcct	ttttcagcta	ttaatgattt	attgttcata	tgaaacactc	cttttatatt	
17361	aatttgatat	tgataccacc	aatcaaatgt	gattggtagc	attgtattaa	attaatattc	tggataattt	
17431	attgagaaag	tccagttatc	atcaaatgaa	attgttttat	tttcaagtaa	ctttttagcc	tcatccacct	
17501	caaattctaa	atagaggaat	ttactaagtt	tatcctcatc	tctaaaaatt	ttcatacata	ccacgttatt	
17571	tgaataaatt	tctgtgtata	cgatcggttc	attcatgttt	atcatccttt	ctttattaca	tatatagtat	
17641	atcatgtatt	tacatatatg	tcaatcattt	aattcattta	ttttaatgat	ttatttgatt	gtttttttat	
17711						aacaattaaa	ttcatataaa	
17781	tgtagtttgg	ggtcagttac	atttgtgtta	tcaaaaaaag	ataatattct	att		

Table 22

## Phage 182 ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	182ORF001	2	59667780		Tail protein;
2	182ORF002	1	21523873	573	DNA polymerase;
3	182ORF003	11	1130512639	444	
4	182ORF004	3	46265954		Major head protein;
5	182ORF005	3	1265113700	349	Glycyl-Glycine endopeptidase; Lysostaphin precursor; Encapsidation protein; ATG/GTP-binding site motif A;
6	182ORF006	1	1499516026	343	Upper collar protein;
7	182ORF007	1	77958775	326	Upper collar protein;     Lysozyme; Muramidase;
8	182ORF008	2	1410514983	292 281	Terminal protein;
9	182ORF010	2	13102155		Lower collar protein;
10	182ORF009	2	87659601	183	Pre-neck appendage protein;
11	182ORF011	1 1	960710158 1087211294	140	Fie-reck appendage protein,
12	182ORF012	3	1045610860	134	
13	1820RF013 1820RF014	3	1371614108		Lysis protein;
14		2	8541225		Early protein;
15	1820RF015 1820RF018	-2	1642916737	102	Lany plotess,
16 17		3	1015810454		Leucine-zipper motif;
18	182ORF020 182ORF019	3	43234613	96	Head protein;
19	1820RF019	-3	1674917033	94	i rous protein,
20	1820RF022	1	1286813149	93	
21	1820RF023	-2	1191412189	91	
22	182ORF017	1	154426	90	
23	182ORF024	3	61746446	90	· · · · · · · · · · · · · · · · · · ·
24	182ORF025	2	548814	88	Early protein;
25	182ORF026	-3	1299913259	86	
26	182ORF027	-1	1464214896	84	
27	182ORF028	3	1443014672	80	
28	182ORF021	-3	1710617339	77	
29	182ORF030	-1	1619916429	76	
30	182ORF031	-3	83798603	74	
31	1820RF032	-1	1119511413	72	
32	182ORF033	-1	47274942	71	
33	1820RF034	-1	59516160	69	
34	182ORF029	-3	1741217606	64	
35	182ORF035	-3	1557015758	62	
36	182ORF036	-3	21272315	62	
37	182ORF037	-1	1209512280	61	
38	182ORF038	3	1476914951	60	<u> </u>
39	1820RF039	2	999210171	59	
40	182ORF040	-3	1602916202	57	
41	1820RF041	1	38864056	56	Early protein;
42	182ORF042	-3	1067110832	53	
43	182ORF043	-3	1049110652	53	
44	182ORF044	-1	62996457	52	
45	1820RF045	-2	65716729	52	
46	182ORF046	2	23722527	51	
47	182ORF047	•2	1320113353	50	
48	182ORF048	-3	32433395	50 48	
49	182ORF049	3	15781724		
50	182ORF050	2	80128155	46	
51	182ORF051	3	93909530	45	1
52	182ORF052	1	40964233 1565615793	45	
53	182ORF053	-2	80028136	44	
54 55	182ORF054 182ORF055	2	83248455	43	
56	182ORF056	3	65496680	43	
57	182ORF057	-3	81338264	43	
	182ORF058	-3 -1	50485176	42	
58 59	182ORF059	-2	1574815876	42	
60	1820RF060	-3	1527615404	42	-
	182ORF061	-3	19742102	42	
64 !	102UKTUU I				
61	1820PE062	-2	1867 1992	41	
61 62 63	182ORF062 182ORF063	-2 -3	18671992 1418114306	41	

182ORF065	-2	34603582	40
182ORF066	1	42344353	39
182ORF067	-1	1376313882	39
182ORF068	-1	71487267	39
182ORF069	-3	49085027	39
182ORF070	-3	9121031	39
182ORF071	2	1174111857	38
182ORF072	-3	1161011723	37
182ORF073	-3	27632876	37
182ORF074	-1	88138923	36
182ORF075	-3	73537463	36
182ORF076	-3	23162426	36
182ORF077	2	1185811965	35
182ORF078	-2	75647671	35
182ORF079	-2	73817488	35
182ORF080	-2	43724473	33
	182ORF066 182ORF067 182ORF068 182ORF069 182ORF071 182ORF072 182ORF073 182ORF074 182ORF075 182ORF075 182ORF076 182ORF076 182ORF077 182ORF077	182ORF066 1 182ORF067 -1 182ORF068 -1 182ORF069 -3 182ORF070 -3 182ORF071 2 182ORF072 -3 182ORF073 -3 182ORF074 -1 182ORF075 -3 182ORF076 -3 182ORF077 -2 182ORF077 2 182ORF077 -2 182ORF0778 -2 182ORF0778 -2	182ORF066         1         4234.4353           182ORF067         -1         1376313882           182ORF068         -1         7148.7267           182ORF069         -3         49085027           182ORF070         -3         9121031           182ORF071         2         1174111857           182ORF072         -3         1161011723           182ORF073         -3         27632876           182ORF074         -1         88138923           182ORF075         -3         73537463           182ORF076         -3         23162426           182ORF077         2         1185811965           182ORF078         -2         75647671           182ORF079         -2         73817488

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### Table 23

Predicted amino acid sequences of ORFs from phage 182

```
1820RF001
      5966
      MARRYTN V K L L A N V P F D N T Y T H T R W F K T
6050
      caacaggaacaggaatcgtactttaattcgtttcctgttcttaacgagaatagagattgttcttatcaaagggatacacaactc
      Q Q E Q E S Y F N S F P V L N E N R D C S Y Q R D T Q L
29
      gggggagtttttagagtagataaacacaaagacgccttatatgcttgtaactatctcatctttaaaaacgaagaaacttatcct
6134
      G G V F R V D K H K D A L Y A C N Y L I F K N E E T Y P
57
      6218
85
6302
      113
      Q T Y R F D I G I R E S F I A K E H P Q L Y Y S N G I P
      6386
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#### 1820RF010

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                {\tt caacaccacccgatcatattagaggaacatacaatcatgttcatggaaaagaaatcccatcaatggtgtggacacctgaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaacaatggaa
 14853
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 29
 14937
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                LIFT *
 1820RF039
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 9992
 1
                M L L M I E H F G I R F N A T I L I M E P I L L T R Y V
10076
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 57
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 1820RF040
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29
16034
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                K *
57
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3886
                {\tt atggaactatataaaagcaatgtttatcgtacgtgatgaaggtactattgacggttacgatactgaacactatgtagatatttct}
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3970
                ttacatgactttgaagaaatatatggaaaagaaacacgtgaaattgaagcagtaacattagtaaaaaacaggaaatttaaaaaaa
                LHDFEEIYGKETREIEAVTLVKTGNLKK
29
4054
                taa 4056
57
1820RF042
10832
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               10748
                TĀLĪLFĀ V V FĀCSĀ V C CĀ V PĀF V NĀ
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10652
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1
10568
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29
               S A L T N S R L E S P V R T F L A I V T G I K R L
1820RF044
6457
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6373
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1820RF045
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6729
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13269
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1820RF048
3395
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6633
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8180
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5176
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15792
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15320
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7272
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3498
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4318
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1820RF067
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7267
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1820RF069
5027
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4943
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1031
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947
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11741
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2876
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2792
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29
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1820RF074
8923
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8839
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29
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1820RF075
7463
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7379
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1820RF076
2426
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- \_\_\_\_\_

#### Table 24

Sequence similarities phage 182 and public databases

```
Phage: 182
Database: nr
Query= sid|110156|lan|1820RF001 Phage 182 ORF|5966-7780|2
          (604 letters)
gi | 138124 | sp | P07534 | VG9 BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                            384 e-105
gi|138123|sp|P04331|VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                            374 e-103
gi|1429238|gnl|PID|e1173412 (X99260) tail protein [Bacteriophag...
                                                                            346 3e-94
gi|215339 (M12456) p9 tail protein [Bacteriophage phi-29] >gi|2...
gi|1181970|gnl|PID|e221269 (Z47794) tail protein [Bacteriophage...
gi|1181968|gnl|PID|e221267 (Z47794) tail protein [Bacteriophage...
                                                                            208 8e-53
                                                                             62
                                                                                 8e-09
                                                                             56 6e-07
gi 2500030 sp Q59968 CARA_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM...
                                                                             49 8e-05
Query= sid | 110157 | lan | 1820RF002 Phage 182 ORF | 2152-3873 | 1
          (573 letters)
gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0...
gi|1429230|gn1|PID|e1173404 (X99260) DNA polymerase [Bacterioph...
                                                                            657 0.0
gi 118849 sp | P03680 | DPOL_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP...
                                                                            654 0.0
gi|118851|sp|P06950|DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...
                                                                            654 0.0
gi|15732 (X53371) DNA polymerase (AA 1-575) [Bacteriophage phi-29]
                                                                            651 0.0
gi 15734 (X53370) DNA polymerase (AA 1-575) [Bacteriophage phi-29]
                                                                            651 0.0
gi|1572479|gn1|PID|e242301 (X96987) DNA polymerase {Bacteriopha...
                                                                            565
                                                                                 e-160
gi|1072656|pir||S51275 DNA polymerase - phage CP-1 >gi|836593|g...
gi|118847|sp|P22374|DPOM_ASCIM PROBABLE DNA POLYMERASE >gi|8385...
                                                                            301 le-80
                                                                             71 3e-11
gi 461962 sp P33537 DPOM NEUCR PROBABLE DNA POLYMERASE >gi 2833...
                                                                             65 1e-09
gi 461963 sp P33538 DPOM_NEUIN PROBABLE DNA POLYMERASE >gi 1018...
gi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum po...
gi 2435429 (AF012250) unassigned reading frame (possible DNA po...
                                                                             61 3e-08
gi|578157|gnl|PID|e246743 (X52106) DNA polymerase [Neurospora i...
                                                                             59
                                                                                 1e-07
                                                                             58
gi|2147969|pir||S72369 probable DNA-polymerase - Gelasinospora ...
                                                                                 2e-07
gi|2147968|pir||S62752 probable DNA-polymerase - Gelasinospora ...
                                                                             58 2e-07
gi 3511140 (AF061244) B type DNA polymerase (Agrocybe aegerita)
                                                                             57 3e-07
gi | 118850 | sp | P10479 | DPOL BPPRD DNA POLYMERASE (PROTEIN P1) >gi | ...
                                                                             56 6e-07
gi|578144 (X63909) putative DNA-polymerase, B-type [Morchella c... gi|232013|sp|P30322|DPOM_AGABT PROBABLE DNA POLYMERASE >gi|3208...
                                                                             47
                                                                                 3e-04
                                                                             46 6e-04
Query= sid | 110159 | lan | 1820RF004 | Phage 182 ORF | 4626-5954 | 3
          (442 letters)
gi|138117|sp|P13849|VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN ...
gi|138118|sp|P07531|VG8_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN ...
                                                                           305
                                                                                 3e-82
gi|1429236|gnl|PID|e1173410 (X99260) major head protein [Bacter...
                                                                            300 le-80
gi|1181958|gnl|PID|e221257 (Z47794) major head protein [Bacteri...
                                                                           152 6e-36
Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
          (349 letters)
gi|137932|sp|P15132|VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR...
                                                                            52 8e-06
gi|1429242|gn1|PID|e1173416 (X99260) morphogenesis protein [Bac...
                                                                             48
                                                                                 7e-05
gi 137933 sp P07538 VG13_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR...
                                                                             47 2e-04
Query= sid|110161|lan|182ORF006 Phage 182 ORF|14995-16026|1
         (343 letters)
gi|137944|sp|P11014|VG16_BPPH2 ENCAPSIDATION PROTEIN (LATE PROT...
                                                                           402 e-111
gi|137945|sp|P07541|VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROT...
                                                                           402 e-111
gi|1429245|gnl|PID|e1173419 (X99260) encapsidation protein [Bac...
                                                                           381 e-105
gi|1181972|gn1|PID|e221271 (Z47794) encapsidation protein (Bact...
                                                                           159 2e-38
```

(123 letters)

```
gi|1429239|gn1|PID|e1173413 (X99260) upper collar protein (Bact...
                                                                        271 5e-72
 gi|137915|sp|P07535|VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...
                                                                        256 le-67
 gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...
                                                                        256
                                                                             2e-67
 gi|1181960|gn1|PID|e221259 (Z47794) connector protein (Bacterio...
                                                                        148 6e-35
 Query= sid|110163|lan|1820RF008 Phage 182 ORF|14105-14983|2
          (292 letters)
 gi|4210750|gnl|PID|e1374037 (AJ132604) LysL protein [Lactococcu...
                                                                        139 2e-32
 gi 462559 sp P34020 LYC CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC...
                                                                         75
                                                                             86-13
 gi 2327014 (U82823) putative lysozyme (Saccharopolyspora erythr...
                                                                         64
                                                                             2e-09
 gi | 126652 | sp | P25310 | LYCM_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA-...
                                                                         60
                                                                             2e-08
 gi | 127789 | sp | P19386 | LYCA BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE...
                                                                         60
                                                                             2e-08
 gi 67761 pir | MUBPCP N-acetylmuramoyl-L-alanine amidase (EC 3.5...
                                                                         59
                                                                             3e-08
 gi 4105636 (AF049087) lys [Leuconostoc oenos bacteriophage 10MC]
                                                                         59
                                                                             3e-08
 gi 623084 (L02496) muramidase; muramidase [Bacteriophage LL-H]
                                                                         57
                                                                             1e-07
 gi | 127787 | sp | P15057 | LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE...
                                                                         57 2e-07
 gi | 126597 | sp | P00721 | LYCH_CHASP N,O-DIACETYLMURAMIDASE (LYSOZYME...
                                                                             2e-07
 gi|127788|sp|P19385|LYCA_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE...
                                                                         57 2e-07
gi|67762|pir||MUBPC7 N-acetylmuramoyl-L-alanine amidase (EC 3.5...
                                                                         56
                                                                             3e-07
gi 3025168 sp P76421 YEGX ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN...
                                                                             26-06
                                                                         53
gi|4204413 (AF047001) Lys44 [Oenococcus oeni temperate bacterio...
gi|2116978|gnl|PID|d1020940 (D88151) cortical fragment-lytic en...
                                                                         53 3e-06
                                                                         52
                                                                             5e-06
gi|2392844 (AF011378) lysin (Bacteriophage skl)
                                                                         48 8e-05
Query= sid | 110164 | lan | 1820RF009 Phage 182 ORF | 8765-9601 | 2
         (278 letters)
gi|1429240|gn1|PID|e1173414 (X99260) lower collar protein (Bact...
                                                                        180 le-44
gi|137921|sp|P04333|VG11_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE...
                                                                        171
                                                                             5e-42
gi 215341 (M12456) pl1 lower collar protein [Bacteriophage phi-29]
                                                                             9e-20
                                                                         98
gi 224162 prf | 1011232B protein pll, lower collar (Bacteriophage...
                                                                         97 1e-19
gi 535260 (Z30339) STARP antigen [Plasmodium reichenowi]
                                                                         50 1e-05
gi 4049753 (AF063866) ORF MSV230 hypothetical protein (Melanopl...
                                                                         49
                                                                             4e-05
gi|2131557|pir||S70306 hypothetical protein YEL077c - yeast (Sa...
                                                                             5e-05
gi | 131782 | sp | P12753 | RA50 YEAST DNA REPAIR PROTEIN RAD50 (153 KD...
                                                                             7e-05
gi|2131309|pir||S70305 hypothetical protein YBL113c - yeast (Sa...
                                                                         47
                                                                             2e-04
gi 499325 (Z26314) STARP antigen [Plasmodium falciparum]
                                                                             3e-04
                                                                         46
gi 3845171 (AE001391) ribosome releasing factor (OO, TP) [Plasm...
                                                                         46
                                                                             3e-04
gi 731903 sp P40434 YIR7 YEAST HYPOTHETICAL 197.5 KD PROTEIN IN...
                                                                             5e-04
                                                                         45
gi|1632829|gnl|PID|e276379 (Y08924) AARP2 protein (Plasmodium f...
                                                                         45 5e-04
gi 1176490 sp P40889 YJW5_YEAST HYPOTHETICAL 197.6 KD PROTEIN I...
                                                                         45
gi|1077300|pir||S51848 hypothetical protein HRD1054 - yeast (Sa...
                                                                             5e-04
gi|2425143 (AF020407) WimA (Dictyostelium discoideum)
                                                                         45
                                                                             6e-04
gi 1181961 | gnl | PID | e221260 (Z47794) collar protein (Bacteriopha...
                                                                             6e~04
                                                                         45
gi|2132657|pir||S64819 probable membrane protein YLL067c - yeas...
                                                                         45
                                                                             8e-04
gi|2133041|pir||S65341 probable membrane protein YPR204w - yeas...
                                                                         45
                                                                             8e-04
gi 730275 sp P39793 PBPA BACSU PENICILLIN-BINDING PROTEINS 1A/1...
                                                                             8e-04
Query= sid|110165|lan|1820RF010 Phage 182 ORF|1310-2155|2
         (281 letters)
gi|135604|sp|P06812|TERM_BPNF DNA TERMINAL PROTEIN >gi|75815|pi...
                                                                         69 3e-11
gi|1572478|gnl|PID|e242334 (X96987) terminal protein [Bacteriop...
                                                                         65
                                                                            3e-10
gi 1429231 gnl PID e1173405 (X99260) terminal protein (Bacterio...
                                                                         64 le-09
Query= sid|110166|lan|1820RF011 Phage 182 ORF|9607-10158|1
         (183 letters)
gi 137928 sp P07537 VG12_BPPZA PRE-NECK APPENDAGE PROTEIN (LATE...
                                                                         51 6e-06
gi|1429241|gnl|PID|e1173415 (X99260) pre-neck appendage protein...
                                                                         51 6e-06
gi | 137927 | sp | P20345 | VG12 BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE...
                                                                        50 le-05
Query= sid | 110169 | lan | 1820RF014 Phage 182 ORF | 13716-14108 | 3
         (130 letters)
gi|137936|sp|P11188|VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14...
                                                                            6e-20
gi 137938 sp P07539 VG14 BPPZA LYSIS PROTEIN (LATE PROTEIN GP14...
                                                                        96
                                                                            8e-20
gi|1429243|gnl|PID|e1173417 (X99260) lysis protein [Bacteriopha...
                                                                        96
                                                                            8e-20
gi|215332 (M14782) lysis protein (Bacteriophage phi-29)
                                                                        94
                                                                            5e-19
Query= sid|110170|lan|1820RF015 Phage 182 ORF|854-1225|2
```

gi 15670 (V01155) reading frame 10 (may be gene 4) [Bacteriopha gi 138072 sp P06953 VG5A_BPPZA EARLY PROTEIN GP5A >gi 75836 pir		5e-12 7e-12
Query= sid 110174 lan 182ORF019 Phage 182 ORF 4323-4613 3 (96 letters)		
gi 1429235 gnl PID e1173409 (X99260) head morphogenesis protein gi 138111 sp P13848 VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE gi 138112 sp P07533 VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE	57	2e-09 3e-08 1e-07
Query= sid 110180 lan 1820RF025 Phage 182 ORF 548-814 2 (88 letters)		
gi 138099 sp P06955 VG6_BPPZA_EARLY_PROTEIN_GP6 >gi 75841 pir   gi 138098 sp P03685 VG6_BPPH2_EARLY_PROTEIN_GP6 >gi 75840 pir   gi 1429234 gn1 PID e1173408 (X99260) gene 6 product [Bacterioph	54	7e-08 2e-07 2e-07

WO 00/32825 PCT/IB99/02040

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#### Table 25

#### Homologies between 182 ORFs and proteins in public databases

Phage: 182 Database: Swissprot Ouery= sid | 110156 | lan | 1820RF001 Phage 182 ORF | 5966-7780 | 2 (604 letters) gi|138124|sp|P07534|VG9\_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) 384 e-106 gi | 138123 | sp | P04331 | VG9 BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) 374 e-103 gi 2500030 sp Q59968 CARA\_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM... 49 2e-05 Query= sid|110157|lan|1820RF002 Phage 182 ORF|2152-3873|1 (573 letters) gi|118848|sp|P19894|DPOL\_BPM2 DNA POLYMERASE 665 0.0 gi 118849 sp P03680 DPOL\_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP2) 654 0.0 gi|118851|sp|P06950|DPOL\_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2) 654 0.0 gi | 118847 | sp | P22374 | DPOM\_ASCIM PROBABLE DNA POLYMERASE 71 7e-12 gi 461962 sp | P33537 | DPOM\_NEUCR PROBABLE DNA POLYMERASE 65 3e-10 gi 461963 sp P33538 DPOM\_NEUIN PROBABLE DNA POLYMERASE 62 3e-09 gi 118850 sp P10479 DPOL\_BPPRD DNA POLYMERASE (PROTEIN P1) 56 2e-07 gi|232013|sp|P30322|DPOM\_AGABT PROBABLE DNA POLYMERASE 46 2e-04 gi|118887|sp|P10582|DPOM\_MAIZE DNA POLYMERASE (S-1 DNA ORF 3) 46 2e-04 Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3 (442 letters) gi|138117|sp|P13849|VG8\_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN ... gi | 138118 | sp | P07531 | VG8\_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN ... Query= sid|110160|lan|1820RF005 Phage 182 ORF|12651-13700|3 (349 letters) 52 2e-06 gi|137932|sp|P15132|VG13\_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR... gi 137933 sp P07538 VG13\_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR... Query= sid|110161|1an|1820RF006 Phage 182 ORF|14995-16026|1 (343 letters) gi|137945|sp|P07541|VG16\_BPPZA ENCAPSIDATION PROTEIN (LATE PROT... 402 e-112 gi 137944 sp P11014 VG16 BPPH2 ENCAPSIDATION PROTEIN (LATE PROT... 402 e-112 Query= sid|110162|lan|1820RF007 Phage 182 ORF|7795-8775|1 (326 letters) gi|137915|sp|P07535|VG10\_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ... 256 3e-68 gi 137914 sp P04332 VG10 BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ... Query= sid|110163|1an|182ORF008 Phage 182 ORF|14105-14983|2 (292 letters) gi|462559|sp|P34020|LYC\_CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC... 75 2e-13 gi | 126652 | sp | P25310 | LYCM\_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA-... 60 5e-09 gi|127789|sp|P19386|LYCA\_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE... 5e-09 gi 127787 sp P15057 LYCA\_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE... gi 126597 sp P00721 LYCH\_CHASP N,O-DIACETYLMURAMIDASE (LYSOZYME... 4e-08 57 4e-08 gi|127788|sp|P19385|LYCA\_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE... 57 5e-08 gi|3025168|sp|P76421|YEGX\_ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN... 53 Se-07 \_\_\_\_ Query= sid | 110164 | lan | 1820RF009 Phage 182 ORF | 8765-9601 | 2 (278 letters) gi|137921|sp|P04333|VG11\_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE... 171 1e-42 gi|131782|sp|P12753|RA50\_YEAST DNA REPAIR PROTEIN RAD50 (153 KD... 48 2e-05 gi 1176490 sp P40889 YJW5 YEAST HYPOTHETICAL 197.6 KD PROTEIN I... 45 1e-04 gi 731903 | sp | P40434 | YIR7 YEAST HYPOTHETICAL 197.5 KD PROTEIN IN... 45 le-04 gi 730275 sp P39793 PBPA\_BACSU PENICILLIN-BINDING PROTEINS 1A/1... 45 2e-04 gi|1168610|sp|P41696|AZF1\_YEAST ASPARAGINE-RICH ZINC FINGER PRO... 44 3e-04

gi 731587 sp P38900 YH19_YEAST HYPOTHETICAL 70.1 KD PROTEIN IN	44	3e-04
Query= sid 110165 lan 182ORF010 Phage 182 ORF 1310-2155 2 (281 letters)		
gi 135604 sp P06812 TERM_BPNF DNA TERMINAL PROTEIN	69	8e-12
Query= sid 110166 lan 182ORF011 Phage 182 ORF 9607-10158 1 (183 letters)		
gi 137928 sp P07537 VG12_BPPZA PRE-NECK APPENDAGE PROTEIN (LATE gi 137927 sp P20345 VG12_BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE	51 50	2e-06 3e-06
Query= sid 110169 1an 1820RF014 Phage 182 ORF 13716-14108 3 (130 letters)		
gi 137936 sp P11188 VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)	97	2e-20
gi 137938 sp P07539 VG14_BPPZA LYSIS PROTEIN (LATE PROTEIN GP14)	96	2e-20
Query= sid 110170 lan 1820RF015 Phage 182 ORF 854-1225 2 (123 letters)		
gi 138072 sp P06953 VG5A_BPPZA EARLY PROTEIN GP5A	69	2e-12
Query= sid 110174 lan 1820RF019 Phage 182 ORF 4323-4613 3 (96 letters)		
gi 138111 sp P13848 VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE gi 138112 sp P07533 VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE	57 54	9e-09 4e-08
Query= sid 110180 lan 1820RF025 Phage 182 ORF 548-814 2 (88 letters)		
gi 138099 sp P06955 VG6_BPPZA EARLY PROTEIN GP6 gi 138098 sp P03685 VG6_BPPH2 EARLY PROTEIN GP6	55 54	2e-08 5e-08

- -

BLASTP 2.0.8 (Jan-05-1999)

Query= sid|110156|lan|182ORF001 Phage 182 ORF|5966-7780|2 (604 letters)

>gi | 138124 | sp | P07534 | VG9\_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >gi|75849|pir||WMBP9Z gene 9 protein - phage PZA >gi|216058 (M11813) tail protein (Bacteriophage PZA) Length = 599

Score = 384 bits (975), Expect = e-105 Identities = 231/610 (37%), Positives = 344/610 (55%), Gaps = 36/610 (5%)

TNVKLLANVPFDNTYTHTRWFKTQQEQESYFNSFPVLNENRDCSYQRDTQLGGVFRVDKH 65 TNV++LA+VPF N Y +TRWF + Q ++FNS + E ++Q + V

TNVRILADVPFSNDYKNTRWFTSSSNQYNWFNSKTRVYEMSKVTFQGFRENKSYISVSLR 68

Query: 66 KDALYACNYLIFKNEETYPSKWQYAFVTDIEYKNDNTSFVTFEIDVLQTYRFDIGIRESF 125 D LY +Y++F+N + Y +KW YAFVT++EYKN T++V FEIDVLQT+ F+I +ESF

Sbjct: 69 LDLLYNASYIMFQNAD-YGNKWFYAFVTELEYKNVGTTYVHFEIDVLQTWMFNIKFQESF 127

Query: 126 IAKEHPQLYYSNGIPFINTIEESLDYGREYTTTNVTTFHPNDGVNFLVILTSEAM--PVG 183 I +BH +L+ +G P INTI+E L+YG EY +V P D + FLV+++ M G

Sbjct: 128 IVREHVKLWNDDGTPTINTIDEGLNYGSEYDIVSVENHRPYDDMMFLVVISKSIMHGTAG 187

Query: 184 DKEDKSG---GSIVGGPSPFSYYLLPINSSGEVYKPN-GAGNANFGEYMAFLT---TKEP 236 S+ G P P YY+ P G+V K G NAN + E +

Sbjct: 188 EAESRLNDINASLNGMPQPLCYYIHPPYKDGKVPKTFIGDNNANLSPIVNMLTNIFSQKS 247

Query: 237 FLNKIVGMYVTSYTGIPFIVDHANKTVRYNAGGSYKIMLPTYASDPTGTMKTFAFFCVKE 296

+N IV MYVT Y G+ + +K ++ + + + A D G + T VK+
Sbjct: 248 AVNNIVNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGI---ADDKHGNVDTIF---VKK 301

Query: 297 ARTFVPKRIDLVGNVYNYFREAFPFNVKESKLFMYPYCLIEITDTKGHVMTLRPEYLTGG 356

+ ID G+ + F + +ESKL MYPYC+ E+TD KG+ M L+ EY+
Sbjct: 302 IPDYETLEID-TGDKWGGFTKD----QESKLMMYPYCVTEVTDFKGNHMNLKTEYIDNN 355

Query: 357 KLSVYVKGSLGISNKVMIEPIDYDVSNSTI----ITNLSDKMLIDNDPNDVGVKSDYASA 412 KL + V+GSLG+SNKV DY+ S +T D LI+N+PND+ + +DY SA

Sbjct: 356 KLKIQVRGSLGVSNKVAYSIQDYNAGGSLSGGDRLTASLDTSLINNNPNDIAIINDYLSA 415

Query: 413 FMQGNKNSLIAQEQNIRNTFRHGMGNSAMSTGGAIFSALASNNPFVGLTNIMGAGQQVNN 472 ++QGNKNSL Q+ +I GM +S G ++ +PF +++ G N Sbjct: 416 YLQGNKNSLENQKSSILFNGIVGMLGGGVSAG----ASAVGRSPFGLASSVTGMTSTAGN 471

Query: 473 YVSEKENGLNLLAGKVADIENIPDNVTQLGSNLSFTTGN-FQNYYQLRFKQIKYEYATRL 531

V + + L K ADI NIP +T++G N +F GN ++ Y ++ KQ+K EY L Sbjct: 472 AVLD----MQALQAKQADIANIPPQLTKMGGNTAFDYGNGYRGVYVIK-KQLKAEYRRSL 526

Query: 532 DRYFSMYGTKSNRVATPNLQTRKAWNFIKLKEPNIVGTMSNDVLTRVKQIFSAGVTLWHT 591

+F YG K NRV PNL+TRKA+N+I+ K+ I G ++N+ L ++ IF G+TLWHT

Sbjct: 527 SSFFHKYGYKINRVKKPNLRTRKAYNYIQTKDCFISGDINNNDLQEIRTIFDNGITLWHT 586

Query: 592 NDVLNYNQDN 601

+D+ NY+ +N

Sbjct: 587 DDIGNYSVEN 596

Query= sid | 110157 | lan | 1820RF002 Phage 182 ORF | 2152-3873 | 1 (573 letters)

>gi|118848|sp|P19894|DPOL BPM2 DNA POLYMERASE >gi|76896|pir||JQ0161 DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2 >gi|215509 (M33144) DNA polymerase (Bacteriophage M2) Length = 572

Score = 665 bits (1697), Expect = 0.0 Identities = 327/589 (55%), Positives = 420/589 (70%), Gaps = 38/589 (6%)

KKYTGDFETTTDLNDCRVWSWGVCDIDNVDNMTFGLEIDSFFEWCKMQGSTDIYFHNEKF 62 K ++ DFETTT L+DCRVW++G +I N+DN G +D F +W M+ D+YFHN KF

Sujec:	4	WESCHIETTIKENCKANNIGINETGUNDMIKIGNSENELMÄNA-METÄMPELUNDVE 95							
Query:	63	DGEFMLSWLFKNGFKWCKEAKEDRTFSTLISNMGQWYALEICWEVNYXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX							
Sbjct:	63	DGAFIVNWLEQHGFKWSNEGLPN-TYNTIISKMGQWYMIDICFGYKGKRKL 112							
Query:	123	XXIIYDSLKKYPFPVKQIAEAFNFPIKKGEIDYTKERPIGYKPTKDEWEYLKNDIQIMAM 182 +IYDSLKK PFPVK+IA+ P P+ KG+IDY ERP+G++ T +E+EY+KNDI+I+A							
Sbjct:	113	HTVIYDSLKKLPFPVKKIAKDFQLPLLKGDIDYHTERPVGHEITPEEYEYIKNDIEIIAR 172							
Query:	183	ALKIQFDQGLTRMTRGSDALGDYKDWLKATHGKSTFKQWFPILSLGFDKDLRKAYKGGFT 242 AL IQF QGL RMT GSD+L +KD L F + FP LSL DK++RKAY+GGFT							
Sbjct:	173	AL IQF QGL RMT GSD+L +KD L F + FF LSL DK++RRAY+GGFT 3 ALDIQFKQGLDRMTAGSDSLKGFKDILSTKKFNKVFPKLSLPMDKEIRKAYRGGFT 228							
Query:	243	WVNKVFQGKEIGDGIVFDVNSLYPSQMYVRPLPYGTPLFYEGEYKPNNDYPLYIQNIKVR 302 W+N ++ KEIG+G+VFDVNSLYPSQMY RPLPYG P+ ++G+Y+ + YPLYIQ I+							
Sbjct:	229	WLNDKYKEKEIGEGMVFDVNSLYPSQMYSRPLPYGAPIVFQGKYEKDEQYPLYIQRIRFE 288							
Query:	303	FRLKEGYIPTIQVKQSSLFIQNEYLESSVNKLGVDELIDLTLTNVDLELFFEHYDILEIH 362 F LKEGYIPTIQ+K++ F NEYL++S GV E ++L LTNVDLEL EHY++ +							
Sbjct:	289	FELKEGYIPTIQIKKNPFFKGNEYLKNSGV-EPVELYLTNVDLELIQEHYELYNVE 343							
Query:	363	YTYGYMFKASCDMFKGWIDKWIEVKNTTEGARKANAKGMLNSLYGKFGTNPDITGKVPYM 422 Y G+ F+ +FK +IDKW VK EGA+K AK MLNSLYGKF +NPD+TGKVPY+							
Sbjct:	344	YIDGFKFREKTGLFKDFIDKWTYVKTHEEGAKKQLAKLMLNSLYGKFASNPDVTGKVPYL 403							
Query:	423	GEDGIVRLTLGEEELRDPVYVPLASFVTAWGRYTTITTAQKCFDRIIYCDTDSIHLVGTE 482 +DG + +G+EE +DPVY P+ F+TAW R+TTIT AQ C+DRIIYCDTDSIHL GTE							
Sbjct:	404	KDDGSLGFRVGDEEYKDPVYTPMGVFITAWARFTTITAAQACYDRIIYCDTDSIHLTGTE 463							
Query:	483	VPEAIDHLVDPKKLGYWGHESTFQRAKFIRQKTYVEEIDGEL 524 VPE I +VDPKKLGYW HESTF+RAK++RQKT YV+E+DG+L							
Sbjct:	464	VPEIIKDIVDPKKLGYWAHESTFKRAKYLRQKTYIQDIYVKEVDGKLKECSPDEATTTKF 523							
Query:	525	NVKCAGMPDRIKEIVTFDNFEVGFSSYGKLLPKRTQGGVVLVDTMFTIK 573 +VKCAGM D IK+ VTFDNF VGFSS GK P + GGVVLVD++FTIK							
Sbjct:	524	SVKCAGMTDTIKKKVTFDNFAVGFSSMGKPKPVQVNGGVVLVDSVFTIK 572							
Query=		110159 lan 1820RF004 Phage 182 ORF 4626-5954 3 442 letters)							
>gi 13	8117	sp P13849 VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN GP8) >gi 75845 pir  WMBP89 gene 8 protein - phage phi-29 >gi 215325 (M14782) major head protein [Bacteriophage phi-29] >gi 225362 prf  1301270B gene 8 [Bacillus sp.] Length = 448							
		309 bits (783), Expect = 2e-83 s = 176/440 (40%), Positives = 250/440 (56%), Gaps = 27/440 (6%)							
Query:	4	KITEQDVLRATNVETPVQLMTAIYNSSSSLFQANVPMPNADNIEAVGAGITRLDVVKNEF 63							
Sbjct:	2	+IT DV + + ++ AI NS F++ VP+ A+N+ VGAGI V+N+F RITFNDVKTSLGITESYDIVNAIRNSQGDNFKSYVPLATANNVAEVGAGILINQTVQNDF 61							
Query:		ISTLVDRIGKVVIRYKSWRNPLKMFKKGNMPLGRTIEEIFVDIAQEHKFNPDESVTGVFK 123							
Sbjct:		I++LVDRIG VVIR S NPLK FKKG +PLGRTIEEI+ DI +E +++ +E+ VF+ ITSLVDRIGLVVIRQVSLNNPLKKFKKGQIPLGRTIEEIYTDITKEKQYDAEEAEQKVFE 121							
Query:	124	QEVPDVKTLFHEINREGYYKQTIQEAWLEKAFTSWDNFNSFVAGVMNALYTGDEVSEFEY 183							
Sbjct:	122	+E+P+VKTLFHE NR+G+Y QTIQ+ L+ AF SW NF SFV+ ++NA+Y EV E+EY REMPNVKTLFHERNRQGFYHQTIQDDSLKTAFVSWGNFESFVSSIINAIYNSAEVDEYEY 181							
Query:	184	TKLLIANYQEKELFKEIBIGEITESNAKEFIRKIKSTSNKLEFMSSAYNAQGVKTS 239							
Sbjct:	182	KLL+ NY K LF ++I E T S EF++K+++T+ KL S +N+ V+T 2 MKLLVDNYYSKGLFTTVKIDEPTSSTGALTEFVKKMRATARKLTLPQGSRDWNSMAVRTR 241							
Query:	240	TSKSDQYXXXXXXXXXXXXXXXFNMSKTDFVGHKIVIDEFPKKEGEESSNIVAVIV 299 + D + FNM++TDF+G+ VID F S+ + AV+V							
Sbjct:	242	SYMEDLHLIIDADLEAELDVDVLAKAFNMNRTDFLGNVTVIDGFASTGLEAVLV 295							
Query:	300	DSEWFMIYDKLYKTTSLYNPEGLYWNYWLHHHQLYSTSQFGNAVAFVKSATKPVTKVAFA 359							
Sbjct:	296	D +WFM+YD L+K ++ NP GLYWNY+ H Q S S+F NAVAFV VT+V + DKDWFMVYDNLHKMETVRNPRGLYWNYYYHVWQTLSVSRFANAVAFVSGDVPAVTQVIVS 355							
		CATTENTIFIC COUNTAI TETRIFEATMOCCEUTICCA DAI UVATUVOTACVATANTIFICI EUC. 410							

```
V ATN + V
                                                        G +T
               +V +G +
Sbjct: 356 PNIAAVKQGGQQQFT---AYVRATNAKDHKV-------VWSVEGGSTGTAI----TG 398
Query: 420 QSLVTFTAIGGQQATVLVTV 439
            L++ + O TV TV
Sbjct: 399 DGLLSVSGNEDNQLTVKATV 418
Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
         (349 letters)
>gi|137932|sp|P15132|VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE
           PROTEIN GP13) >gi|75858|pir||WMBP23 gene 13 protein -
          phage phi-29 >gi|215331 (M14782) morphogenesis protein
           [Bacteriophage phi-29] >gi|225368|prf||1301270H gene 13
           [Bacteriophage phi-29]
          Length = 365
 Score = 51.5 bits (121), Expect = 8e-06
 Identities = 44/166 (26%), Positives = 70/166 (41%), Gaps = 14/166 (8%)
Query: 6 NEQIARGQTIAKILSKYGYNKNSQVGVVANLHWESA---GLNPNSNEXXXXXXXXXX - QWT 61
                 Q I LS G+ K + G++ N+ ES GL N +E
Sbjct: 12 SEMKVNAQYILNYLSSNGWTKQAICGMLGNMQSESTINPGLWQNLDEGNTSLGFGLVQWT 71
Query: 62 PKSNLYRQAQICGLSNAKAETLEGQAEIIAQGDKTGQWMDNTPVSSAGYTNPQTLSAFKQ 121
                                    II + + QW++ ++ Y
                 A GL ++
Sbjct: 72 PASNYINWANSQGLPYKDMDS--ELKRIIWEVNNNAQWINLRDMTFKEY-----IKS 121
Query: 122 SANIDVATINFMCHWERPGKLHIEERLDLAQAYSKHIDGSGGGGVK 167
                  + F+ +ERP + ER D A+ + K++ G GGGG++
Sbjct: 122 TKTPRELAMIFLASYERPANPNQPERGDQAEYWYKNLSGGGGGGLQ 167
Query= sid | 110161 | lan | 1820RF006 Phage 182 ORF | 14995-16026 | 1
         (343 letters)
>gi|137945|sp|P07541|VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROTEIN
          GP16) >gi | 75861 | pir | | WMBP16 gene 16 protein - phage PZA
           >gi|216065 (M11813) morphogenesis protein C
           [Bacteriophage PZA]
          Length = 332
 Score = 402 bits (1023). Expect = e-111
 Identities = 186/332 (56%), Positives = 244/332 (73%), Gaps = 2/332 (0%)
Query: 11 EKNLYYNPNNALGFNCLMLFVIGARGIGKTYGYKKFVVNRFIKHGEQFIYLRRFKTELKK 70
          +K+L+YNP L ++ ++ FVIGARGIGK+Y K + +NRFIK+GEQFIY+RR+K EL K
          DKSLFYNPQKMLSYDRILNFVIGARGIGKSYAMKVYPINRFIKYGEQFIYVRRYKPELAK 61
Sbjct: 2
Query: 71 IPQFFKTMAKEFPDHKLEVKGKEFYCDDKLMGWAVPLSTWGIEKSNEYPEVRTILFDEFL 130
           +F +A+EFPDH+L VKG+ FY D KL GWA+PLS W EKSN YP V TI+FDEF+
Sbjct: 62 VSNYFNDVAQEFPDHELVVKGRRFYIDGKLAGWAIPLSVWQSEKSNAYPNVSTIVFDEFI 121
Query: 131 IEKSKITYLPNEAEALLNMMETVFRRRTNTRCVMLSNATSVVNPYFLYFNLQPDLNKRFN 190
                Y+PNE ALLN+M+TVFR R RC+ LSNA SVVNPYFL+FNL PD+NKRFN
Sbjct: 122 REKDNSNYIPNEVSALLNLMDTVFRNRERVRCICLSNAVSVVNPYFLFFNLVPDVNKRFN 181
Query: 191 LYQDRGILIELCDSKDFAEVKRETPFGRLIRGTEYEDFSINNEFVNDSDTFIEKRSKNSS 250
           +Y D LIE+ DS DF+ +R+T FGRLI GTEY + S++N+F+ DS FIEKRSK+S
Sbjct: 182 VYDD--ALIEIPDSLDFSSERRKTRFGRLIDGTEYGEMSLDNQFIGDSHVFIEKRSKDSK 239
Query: 251 FLCAIAFEGKIFGYWIDAETGCVYVSYDYQPNTNHFYAMTTKDHEENRLLMKWWRNNYYL 310
          F+ +I + G G W+D G +YV + P+T + Y +TT D EN +L+ N++NNY+L
Sbjct: 240 FVFSIVYNGFTLGVWVDVNQGLMYVDTAHDPSTKNVYTLTTDDLNENMMLITNYKNNYHL 299
Query: 311 STVAKAFKNSYLRFDNIVIKNLHYDLFNKMKI 342
            +A AF N YLRFDN VI+N+ Y+LF KM+I
                                                                             - -
Sbjct: 300 RKLASAFMNGYLRFDNQVIRNIAYELFRKMRI 331
Query- sid | 110162 | lan | 1820RF007 Phage 182 ORF | 7795-8775 | 1
        (326 letters)
>gi|1429239|emb|CAA67658| (X99260) upper collar protein
```

[Bacteriophage B103]

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Length = 308
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Score = 271 bits (685), Expect = 6e-72
 Identities = 131/275 (47%), Positives = 187/275 (67%), Gaps = 5/275 (1%)
Query: 36 YYEHYRRQLTLLTFQLFEWENLPKSIDPRYLEIALHTNGYLGFFKDPTLGFMVCAGAEDG 95
           +Y HY + L L +QLFEWE LP S+DP YLE ++H GY+GF+KDP +G++ C GA G
Sbjct: 22 WYYHYYQYLCSLAYQLFEWERLPPSVDPSYLEKSIHQFGYVGFYKDPRIGYIACQGALSG 81
Query: 96 QIDHYHNPIFFTANEAMYHKRYPVLRYDDDDDKSKCIMLYNNDLKVPTLPSLHRFALDMA 155
            +DHY+ P F A+ Y + + Y D +K+ + +YNNDLK TLP+L FA D+A
Sbjct: 82 TVDHYNLPDRFHASSVGYQNTFKLYNYSDMKEKNMGVAIYNNDLKCSTLPALEMFAQDLA 141
Query: 156 DINQISRVNRRAQKTPVIIQTDEKKYFSLLQAYNQIDENNQAVFVDKDMEFDESFNVWQT 215
           ++ +I VN+ AQKTPV+I ++ SL YNQ + N +FV + ++ D + V++T
Sbjct: 142 ELKEIIAVNQNAQKTPVLIAANDNNQLSLKNIYNQYEGNAPVIFVHESLDLD-NLKVFKT 200
Query: 216 NAPYVVDKLRSELNEVWNEVLTFLGINNANVDKTARVQTSEVLSNNEQIESSGNILLKSR 275
           +APYVVDKL ++ N VWNEV+T+LGI NAN++K R+ TSEV SN+EQIESSGNI LK+R
Sbjct: 201 DAPYVVDKLNAQKNAVWNEVMTYLGIKNANLEKKERMVTSEVDSNDEQIESSGNIYLKAR 260
Query: 276 KEFCDRVNRVFGDELDGKIDVKFRTDAVRQLQLAA 310
           +E C++++ ++G L VKFR D V Q++L A
Sbjct: 261 QEACNKISELYGLNL----KVKFRYDIVEQMRLNA 291
Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
         (292 letters)
>gi|4210750|emb|CAA10710| (AJ132604) LysL protein [Lactococcus
           lactisl
           Length = 235
 Score = 139 bits (347), Expect = 2e-32
 Identities = 85/210 (40%), Positives = 114/210 (53%), Gaps = 14/210 (6%)
Query: 2 MNGIDISSYQTGIDLSKVPCDFVNIKATGGTGYVNPDCDRAFQQALSLGKKIGVYHFAHE 61
           MNGIDISSYQ ++ VP DFV IKAT GT Y+NP + Q + K +G YHFA
          MNGIDISSYQAELNAGIVPSDFVIIKATEGTNYINPTWEEQAGQVIQTNKLLGFYHFAS- 59
Sbict: 1
Query: 62 RGLEGTPQQEAQFFLDNIKGYIGKAVLILDFEGS--NQKDVNWAKAFLDYVYNKTGVKAW 119
              G P EA FF+ +K YIGKAVL+LDFE N
                                                    A+ FL+ V KTG+
Sbict: 60 --- VGNPIAEADFFISVVKNYIGKAVLVLDFEAGAINAWGNVGARQFLNRVKEKTGINPM 116
Query: 120 FYTYTANLNTTDFSSIAKGDYGLWVAEYGSNQPQGYSQPAPPKTNN----FPIVACFQF 174
                     ++S+I+ + LWVA+Y S P GY + P T+
Sbjct: 117 IYMSSDVTRQFNWSTISSTN-PLWVAQYASMNPTGYQ--SEPWTDGKGYGAWSSAAIHOY 173
Query: 175 TSKGRLPGYNGNLDLNVFYGDGNTWDLYVG 204
           +S G L ++GNLD+N+ Y + N W
Sbjct: 174 SSAGSLSNWSGNLDINLAYINANQWKSLAG 203
Query= sid|110164|lan|1820RF009 Phage 182 ORF|8765-9601|2
         (278 letters)
>gi|1429240|emb|CAA67659| (X99260) lower collar protein
           [Bacteriophage B103]
           Length = 293
 Score = 180 bits (451), Expect = 1e-44
 Identities = 115/296 (38%), Positives = 161/296 (53%), Gaps = 33/296 (11%)
          LKRYIESFTYYQPELSRKERIEVGRKQLFDFDYPFYDETKRAEFETKFINHFYLREIGSE 62
Query: 3
          L YIE ++ Y+ LS E+IE GR +LFDF YP +DE+ R FET FI +FY+REIG E
                                                                                  _____
          LSTYIEMWSQYETGLSMAEKIEKGRPKLFDFQYPIFDESYRKVFETHFIRNFYMREIGFE 67
Sbict: 8
Query: 63 TMGSFKFNLDEYLNLNMPYWNKMFLSNLEEF-PIFDDMDYTIDEKQKLLNEIDTNIKANR 121
           T G FKFNL+ +L +NMPY+NK+F S L ++ P+ + T K+
                                                             DT
Sbjct: 68 TEGLFKFNLETWLIINMPYFNKLFESELIKYDPLENTRLNTTGNKKN-----DTERNDNR 122
Query: 122 D-----ESKNQTKQVDQTDNRNKNTRDTGTT----DSFSRNTYTDTPQKDLRIASNG 169 D + K+ TK D+T+ + D TT D+F+R +D P L + +N
```

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Sbjct: 123 DTTGSMKADGKSNTKTSDKTNATGSSKEDGKTTGSVTDDNFNRKIDSDQPDSRLNLTTN- 181
Query: 170 DGTGVINYATNITEDLSKETTSSTGVETNNDKTNQNTRSNAS------EKETKNTD 219
DG G + YA+ I E+ + ++TG TNN ++ + S S T N

Sbjct: 182 DGQGTLEYASAIEENNTNNKRNTTG--TNNVTSSAESESTGSGTSDTVTTDNANTTTNDK 239
Query: 220 INKDQNQTKDTITRYKGKKGNTDYADLLEKYRRSVLRIEKMIFREMNKEGLFLLVY 275
           +N N +D I GK G YA L++ YR ++LRIEK IF EM + LF+LVY
Sbjct: 240 LNSQINNVEDYIESKIGKSGTQSYASLVQDYRAALLRIEKRIFDEMQE--LFMLVY 293
Query= sid | 110165 | lan | 1820RF010 Phage 182 ORF | 1310-2155 | 2
          (281 letters)
>gi|135604|sp|P06812|TERM_BPNF DNA TERMINAL PROTEIN
           >qi|75815|pir||ERBPNP terminal protein - phage NF
           >gi|579177|emb|CAA68440| (Y00363) gene E product (AA
           1-267) [Bacteriophage NF]
           Length = 266
 Score = 74.9 bits (181), Expect = 6e-13
 Identities = 73/275 (26%), Positives = 129/275 (46%), Gaps = 37/275 (13%)
           VRISKNDRAKLEKIYGKSNKARKKYNRLRQK-GVE---ERQLPTVPTSKKRLIDYVKSTN 58
            +RI+ ND+A K+ K+ KA K +R ++K G++ E +LP + + +
           IRITNNDKALYAKLV-KNTKA--KISRTKKKYGIDLSNEIELPPLESFQ------ 52
Sbjct: 7
Query: 59 MSRSDFNKMLDELVDFAQPYNENYIFEINKRNVAISRAQIKEAQIKTEQAQKAKEEHYKE 118
            +R +FNK + F N+NY F NK + S+A+I E T++AQ+ +E +E
Sbjct: 53 -TREEFNKWKQKQESFTNRANQNYQFVKNKYGIVASKAKINEIAKNTKEAQRIVDEQREE 111
Query: 119 L------NKVEVKKPTENTIVTPTILTELGADLPFQAIPDFNIDAFTSPEGVQSYLEN 170
+ K + I++P+ +T G P DFN D S +++ E
Sbjct: 112 IEDKPFISGGKQQGTVGQRMQILSPSQVT--GISRP----SDFNFDDVRSYARLRTLEEG 165
Query: 171 IG-KQDEQYFDERDQLYYDNFRQAMFTIFNSD--ADDIVRLLDSMGLDLFMKTYVSNFLD 227
+ K Y+D R + NF + + FNSD +D++V L + D F + Y+ F +
Sbjct: 166 MAEKASPDYYDRRMTQMHQNFIEIVEKSFNSDWLSDELVERLKKIPPDDFFELYLM-FDE 224
Query: 228 MNLDYIYDEAEVQQKKEQVYSKIAKVIESETGGEV 262
           ++ +Y E E + B + +KI ++ G+V
Sbjct: 225 ISFEYFDSEGEDVEASEAMLNKIHSYLDRYERGDV 259
Query= sid|110166|lan|1820RF011 Phage 182 ORF|9607-10158|1
         (183 letters)
>gi|1429241|emb|CAA67660| (X99260) pre-neck appendage protein
            [Bacteriophage B103]
           Length = 860
 Score = 50.8 bits (119), Expect = 6e-06
 Identities = 29/105 (27%), Positives = 56/105 (52%), Gaps = 6/105 (5%)
           KRFDGLPAVFKERFSKYPHTEYRYELLLDEEVSALIAYLNEVGALVNDMSGYLNYFIEHF 67
           +RF+ L + + + +Y T + + L E+++ +I YLN++G L ND+
           RRFEKLGEMMVQVYERYLPTAFDESMTLLEKMNKIIEYLNQIGRLTNDVVEEWNKVMEWI 66
Sbict: 7
Query: 68 V-EKLEBITNDTLKKWLSDGTLENLINDTVFANYIKBIKRLQILV 111
            + + LE+ +TL+KW +G +L+ I E+K+ + V
Sbjct: 67 LNDGLEDYVKETLEKWYEEGKFADLV----IQVIDELKQFGVSV 106
                                                                                   - -
Query= sid|110169|lan|1820RF014 Phage 182 ORF|13716-14108|3
         (130 letters)
>gi|137936|sp|P11188|VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)
           >gi|75860|pir||WMBP29 gene 14 protein - phage phi-29
           >gi|15678|emb|CAA28631| (X04962) gene 14 product (AA
```

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1-393) [Bacteriophage phi-29] >gi|225369|prf||1301270J
           gene 14 [Bacteriophage phi-29]
           Length = 131
 Score = 96.7 bits (237), Expect = 6e-20
 Identities = 53/131 (40%), Positives = 81/131 (61%), Gaps = 3/131 (2%)
          MIEYITQWL-ADDNHLVYGLIIWLMVAMIIDFVLGFTIAKFNKBIDFSSFKAKAGIIVKV 59
           MI ++ +L D+ L+Y L +LMV M++D VLG AK N I FSSFK K G+++KV
          MIAWMOHPLETDETKLIYWLT-FLMVCMVVDTVLGVLFAKLNPNIKFSSFKIKTGVLIKV 61
Sbict: 3
Query: 60 AEMVLVVYFIPVAVKFGAVGITMYITMLVGLILSEIYSILGHISDIDDDNNWTDYVKKFL 119
           +EM+L + IP AV F A G+ + T+ L +SEIYSI GH+ +DD +++ + ++ F
Sbjct: 62 SEMILALLAIPFAVPFPA-GLPLLYTVYTALCVSEIYSIFGHLRLVDDKSDFLEILENFF 120
Query: 120 DGTLNRKDDIK 130
            T + + K
Sbjct: 121 KRTSGKNKEEK 131
Query= sid|110170|lan|1820RF015 Phage 182 ORF|854-1225|2
         (123 letters)
>gi|15670|emb|CAA24483| (V01155) reading frame 10 (may be gene 4)
           [Bacteriophage phi-29]
           Length = 124
 Score = 69.9 bits (168), Expect = 6e-12
 Identities = 39/119 (32%), Positives = 64/119 (53%), Gaps = 3/119 (2%)
          IVKSTFDTQTPEGMLQVFNATNGASIPLRNAI-GEVLELKDILVYSDEVSGFGGAEPSQA 61
Query: 3
           IVK+TFDT+T EG +++FNA G +N G ++E I Y
          IVKATFDTETLEGQIKIFNAQTGGGQSFKNLPDGTIIEANAIAQYKQVSDTYGDAK--EE 63
Sbict: 6
Query: 62 ELVAFFTEDGKTYAGVSAVATKSAKNLIDMMTANPDIKPKISFVEGKSNGGQKFVNLQV 120
           + F DG Y+ +S ++A +LID++T + K+ V+G S+ G F +LQ+
Sbjct: 64 TVTTIFAADGSLYSAISKTVAEAASDLIDLVTRHKLETFKVKVVQGTSSKGNVFFSLQL 122
Query= sid | 110174 | 1an | 1820RF019 Phage 182 ORF | 4323-4613 | 3
         (96 letters)
>gi|1429235|emb|CAA67654| (X99260) head morphogenesis protein
          {Bacteriophage B103}
         Length = 101
 Score = 60.9 bits (145), Expect = 1e-09
 Identities = 34/96 (35%), Positives = 53/96 (54%), Gaps = 5/96 (5%)
Query: 1 MEIKEHESILNGILESVTDGEARSKIVEHLEALREDYGATTEALTSANSTLEKLKKDNEA 60
         MB HE ILN + + + R+++ L+ LR DYG+ + S EKL+ +N
Sbjct: 3 MERDSHEEILNKLNDPELEHSERTEL---LQQLRADYGSVLSEFSELTSATEKLRAENSD 59
Query: 61 LVISNSKLFRERAIVEPAEN--NEPETDQNITLDDL 94
         L++SNSKLFR+ I + E + E + IT++DL
Sbjct: 60 LIVSNSKLFRQVGITKEKEEEIKQEELSETITIEDL 95
Query= sid|110180|lan|182ORF025 Phage 182 ORF|548-814|2
         (88 letters)
                                                                            _____
>gi|138099|sp|P06955|VG6_BPPZA EARLY PROTEIN GP6
         >gi|75841|pir||ERBP6Z gene 6 protein - phage PZA
         >gi|216047 (M11813) gene 6 product [Bacteriophage PZA]
         >gi|224746|prf||1112171K ORF 6 [Bacteriophage PZA]
         Length = 96
```

Score = 55.0 bits (130), Expect = 8e-08

Identities = 28/79 (35%), Positives = 45/79 (56%)

Query: 4 KLMQRNVTSTKVEFSEVIVQDGAPTIVPCEPVVLTGKLSEEKALSAIKRKNPDKNVVVTN 63 K+MQR +T T V +++++ DG + G LS E+A +KRK + V V + Sbjct: 3 KMMQREITKTTVNVAKMVMVDGEVQVEQLPSETFVGNLSMEQAQWRMKRKYKGEPVQVVS 62

Query: 64 VSHETALYTMPVDKFIELA 82 V T +Y +PV+KF+E+A Sbjct: 63 VEPNTEVYELPVEKFLEVA 81

## Table 26

## Secondary structure prediction for ORF 1820RF008

1	MMNGIDISSY	QTGIDLSKVP	CDFVNIKATG	GTGYVNPDCD	RAFQQALSLG	KKIGVYHFAH
	CCCCCCCCC	CCCCCCCC	CCEEEEECC	cccccccc	нинининис	CCCCEEEEE
61	ERGLEGTPQQ	EAQFFLDNIK	GYIGKAVLIL	DFEGSNQKDV	NWAKAFLDYV	YNKTGVKAWF
	СССССССНН	нинининис	CCCCEEEEE	ССССССННН	нинининин	HCCCCCEEEE
121	YTYTANLNTT	DFSSIAKGDY	GLWVAEYGSN	QPQGYSQPAP	PKTNNFPIVA	CFQFTSKGRL
	EEECCCCCCC	CCCEECCCCC	CEEEEECCCC	cccccccc	CCCCCCEEE	EEEECCCCCC
181	PGYNGNLDLN	VFYGDGNTWD	LYVGKKQDQI	VPPENKIFDA	TSDEFIFTLT	TGSTSVFYFD
	CCCCCCCEE	EEECCCCCCE	EEECCCCCC	cccccccc	CCCEEEEEC	CCCCEEECC
241	GETIFELSDP	TQLDHIRGTY	NHVHGKEIPS	MVWTPEQFDI	YLKMYEKKPV	YK
	CCEEEECCCC	CCHHHHCCEE	CCCCCEECC	ССССССННН	HHHHHCCCCE	EC

## Secondary structure prediction for ORF 182ORF014

- 121 GTLNRKDDIK CCCCCCEEC

#### Table 27

## Enterococcus accession numbers 242/242

gi|2895751|gb|AF044978.1|AF044978 [2895751] gi|4803755|dbj|AB026843.1|AB026843 [4803755] gi|4769001|gb|AF140549.1|AF140549 [4769001] gi|4760901|gb|AF099088.1|AF099088 [4760901] gi|4704705|gb|AF121254.1|AF121254 [4704705] gi|3342117|gb|AF076604.1|AF076604 [3342117] gi|4688824|emb|AJ132470.1|ESP132470 [4688824] gi|4732085|gb|AF125553.1|AF125553 [4732085] gi|4732082|gb|AF125552.1|AF125552 [4732082] gi|4732079|gb|AF125551.1|AF125551 [4732079] gi|4732076|gb|AF125550.1|AF125550 [4732076] gi|4732073|gb|AF125548.1|AF125548 [4732073] gi|4732070|gb|AF125547.1|AF125547 [4732070] gi|4732067|gb|AF125546.1|AF125546 [4732067] gi|4732064|gb|AF125545.1|AF125545 [4732064] gi|4732061|gb|AF125544.1|AF125544 [4732061] gi|4704653|gb|AF114715.1|AF114715 [4704653] gi|4704564|gb|AF102550.1|AF102550 [4704564] gi|4688827|emb|AJ238249.1|EFA238249 [4688827] gi|4680606|gb|AF125198.1|AF125198 [4680606] gi|4633279|gb|AF117609.1|AF117609 [4633279] gi|4633124|gb|AF110130.1|AF110130 [4633124] gi|4590399|gb|AF124258.1|AF124258 [4590399] gi|4590336|gb|AF108380.1|AF108380 [4590336] gi|4590335|gb|AF108379.1|AF108379 [4590335] gi|4019167|gb|U21300.1|CXU21300 [4019167] gi|4545122|gb|AF077816.1|AF077816 [4545122] gi|4433610|gb|AF106614.1|AF106614 [4433610] gi|4468838|emb|AJ132039.1|EFA132039 [4468838] gi|4468121|emb|AJ132958.1|BPH132958 [4468121] gi|4456104|emb|Y17302.1|EHI17302 [4456104] gi|4433611|gb|AF106615.1|AF106615 [4433611] gi|4433607|gb|AF106611.1|AF106611 [4433607]

gi|4098267|gb|U76614.1|BLU76614 [4098267] gi|47019|emb|Y00116.1|SFAMB1 [47019] gi|4158179|emb|AL035206.1|SC9B5 [4158179] gi|4165458|emb|X79343.1|EF16SSPA [4165458] gi|4165457|emb|X79342.1|EFTRNALA [4165457] gi|4165456|emb|X79341.1|EF23SRNA [4165456] gi|4150978|emb|Y14027.1|EFY14027 [4150978] gi|4127803|emb|AJ223161.1|EFAJ3161 [4127803] gi|2956685|emb|Y16413.1|EFENTIJO [2956685] gi|2665346|emb|Y13922.1|EHY13922 [2665346] gi|4324675|gb|AF109375.1|AF109375 [4324675] gi|4234627|gb|AF061013.1|AF061013 [4234627] gi|4234626|gb|AF061012.1|AF061012 [4234626] gi|4234625|gb|AF061011.1|AF061011 [4234625] gi|4234624|gb|AF061010.1|AF061010 [4234624] gi|4234623|gb|AF061009.1|AF061009 [4234623] gi|4234622|gb|AF061008.1|AF061008 [4234622] gi|4234621|gb|AF061007.1|AF061007 [4234621] gi|4234620|gb|AF061006.1|AF061006 [4234620] gi|4234619|gb|AF061005.1|AF061005 [4234619] gi|4234618|gb|AF061004.1|AF061004 [4234618] gi|4234617|gb|AF061003.1|AF061003 [4234617] gi|4234616|gb|AF061002.1|AF061002 [4234616] gi|4234615|gb|AF061001.1|AF061001 [4234615] gi|4234614|gb|AF061000.1|AF061000 [4234614] gi|3138990|gb|AF060241.1|AF060241 [3138990] gi|3138986|gb|AF060240.1|AF060240 [3138986] gi|4204535|gb|AF094803.1|AF094803 [4204535] gi|4204534|gb|AF094802.1|AF094802 [4204534] gi|4204533|gb|AF094801.1|AF094801 [4204533] gi|4204532|gb|AF094800.1|AF094800 [4204532] gi|4204531|gb|AF094799.1|AF094799-[4204531]gi|4204530|gb|AF094798.1|AF094798 [4204530] gi|4204529|gb|AF094797.1|AF094797 [4204529] gi|4204528|gb|AF094796.1|AF094796 [4204528] gi|4204527|gb|AF094795.1|AF094795 [4204527]

gi|4204526|gb|AF094794.1|AF094794 [4204526] gi|4204525|gb|AF094793.1|AF094793 [4204525] gi|4204524|gb|AF094792.1|AF094792 [4204524] gi|4204523|gb|AF094791.1|AF094791 [4204523] gi|4204522|gb|AF094790.1|AF094790 [4204522] gi|4204521|gb|AF094789.1|AF094789 [4204521] gi|4204520|gb|AF094788.1|AF094788 [4204520] gi|4204519|gb|AF094787.1|AF094787 [4204519] gi|4204518|gb|AF094786.1|AF094786 [4204518] gi|4204517|gb|AF094785.1|AF094785 [4204517] gi|4204516|gb|AF094784.1|AF094784 [4204516] gi|4204515|gb|AF094783.1|AF094783 [4204515] gi|4204514|gb|AF094782.1|AF094782 [4204514] gi|4204513|gb|AF094781.1|AF094781 [4204513] gi|4204512|gb|AF094780.1|AF094780 [4204512] gi|3873186|gb|AF034779.1|AF034779 [3873186] gi|4151367|gb|AF093508.1|AF093508 [4151367] gi|2828136|gb|AF039903.1|AF039903 [2828136] gi|2828135|gb|AF039902.1|AF039902 [2828135] gi|2828134|gb|AF039901.1|AF039901 [2828134] gi|2828133|gb|AF039900.1|AF039900 [2828133] gi|2828132|gb|AF039899.1|AF039899 [2828132] gi|2828131|gb|AF039898.1|AF039898 [2828131] gi|4103866|gb|AF028812.1|AF028812 [4103866] gi|4103864|gb|AF028811.1|AF028811 [4103864] gi|2605925|gb|AF029727.1|AF029727 [2605925] gi|1402750|gb|U60038.1|EFU60038 [1402750] gi|1835780|gb|U86375.1|EFU86375 [1835780] gi|3831555|gb|AF047608.1|AF047608 [3831555] gi|3790617|gb|AF097414.1|AF097414 [3790617] gi|3767587|dbj|AB005036.1|AB005036 [3767587] gi|3757810|gb|AF042288.1|AF042288 [3757810] gi|3747039|gb|AF093509.1|AF093509 [3747039] gi|3660559|dbj|AB017811.1|AB017811 [3660559] gi|1147743|gb|U42211.1|EHU42211 [1147743] gi|3676412|gb|AF051917.1|AF051917 [3676412] gi|3676164|emb|AJ011113.1|EFA011113 [3676164] gi|2612869|gb|AF005726.1|AF005726 [2612869] gi|2353762|gb|AF016233.1|AF016233 [2353762]

gi|2149899|gb|U94707.1|EFU94707 [2149899] gi|2149149|gb|U82366.1|LSU82366 [2149149] gi|1469463|gb|U49512.1|EFU49512 [1469463] gi|1244503|gb|U35366.1|EFU35366 [1244503] gi|833854|gb|U26268.1|EFU26268 [833854] gi|841200|gb|U18931.1|CPU18931 [841200] gi|460079|gb|U00457.1|U00457 [460079] gi|460077|gb|U00456.1|U00456 [460077] gi|535661|gb|L34675.1|INSTRANSPO [535661] gi|3023041|gb|AF007787.1|AF007787 [3023041] gi|431124|gb|L15633.1|TRN916ENT [431124] gi|388106|gb|L23802.1|ENEEBSA [388106] gi|3608387|gb|AF071085.1|AF071085 [3608387] gi|3551851|gb|AF076027.1|AF076027 [3551851] gi|3551773|gb|U94770.1|SPU94770 [3551773] gi|3551743|gb|U57498.1|ECU57498 [3551743] gi|3243178|gb|AF063010.1|AF063010 [3243178] gi|3136316|gb|AF063900.1|AF063900 [3136316] gi|3540256|gb|AF052459.1|AF052459 [3540256] gi|755215|gb|U17696.1|LLU17696 [755215] gi|3421437|gb|AF082295.1|AF082295 [3421437] gi|3421436|gb|AF082294.1|AF082294 [3421436] gi|3421435|gb|AF082293.1|AF082293 [3421435] gi|3421434|gb|AF082292.1|AF082292 [3421434] gi|3341430|emb|Y17797.1|EFY17797 [3341430] gi|3319647|emb|X69092.1|EHPBP3RA [3319647] gi|3292886|emb|AJ007584.1|EFA7584 [3292886] gi|3261536|emb|AL021958.1|MTV041 [3261536] gi|3250708|emb|Z95150.1|MTCY164 [3250708] gi|3249688|gb|AF070678.1|AF070678 [3249688] gi|3249687|gb|AF070677.1|AF070677 [3249687] gi|3249686|gb|AF070676.1|AF070676 [3249686] gi|3219158|dbj|AB015233.1|AB015233 [3219158] gi|2765275|emb|Y12924.1|SPY12924 [2765275] gi|3183687|emb|Y11621.1|EA16SRRN [3183687] gi|2765274|emb|Y12923.1|EFY12923 [2765274] gi|2765273|emb|Y12922.1|ESY12922 [2765273] gi|2765272|emb|Y12921.1|ESY12921 [2765272] gi|2765271|emb|Y12920.1|EDY12920 [2765271] gi|2765270|emb|Y12919.1|ESY12919 [2765270]

gi|2765269|emb|Y12918.1|ECY12918 [2765269] gi|2765268|emb|Y12917.1|ECY12917 [2765268] gi|2765267|emb|Y12916.1|EPY12916 [2765267] gi|2765266|emb|Y12915.1|ESY12915 [2765266] gi|2765265|emb|Y12914.1|ERY12914 [2765265] gi|2765264|emb|Y12913.1|EMY12913 [2765264] gi|2765263|emb|Y12912.1|EHY12912 [2765263] gi|2765262|emb|Y12911.1|EMY12911 [2765262] gi|2765261|emb|Y12910.1|EGY12910 [2765261] gi|2765260|emb|Y12909.1|EDY12909 [2765260] gi|2765259|emb|Y12908.1|ECY12908 [2765259] gi|2765258|emb|Y12907.1|EAY12907 [2765258] gi|2765257|emb|Y12906.1|EFY12906 [2765257] gi|2765256|emb|Y12905.1|EFY12905 [2765256] gi|2894541|emb|AJ223332.1|EFAJ3332 [2894541] gi|2894539|emb|AJ223331.1|EFAJ3331 [2894539] gi|3108058|gb|AF060881.1|AF060881 [3108058] gij3087776|emb|AJ223633.1|EFAJ3633 [3087776] gi|3080754|gb|AF016483.1|AF016483 [3080754] gi|2197119|gb|AF003921.1|AF003921 [2197119] gi|2982722|dbj|AB012213.1|AB012213 [2982722] gi|2982721|dbj|AB012212.1|AB012212 [2982721] gi|2058780|gb|B07890.1|B07890 [2058780] gi|2058779|gb|B07889.1|B07889 [2058779] gi|2058778|gb|B07888.1|B07888 [2058778] gi|2058777|gb|B07887.1|B07887 [2058777] gi|2058776|gb|B07886.1|B07886 [2058776] gi|2058775|gb|B07885.1|B07885 [2058775] gi|2058774|gb|B07884.1|B07884 [2058774] gi|2058773|gb|B07873.1|B07873 [2058773] gi|2058772|gb|B07872.1|B07872 [2058772] gi|2058771|gb|B07871.1|B07871 [2058771] gi|2058770|gb|B07870.1|B07870 [2058770] gi|2058769|gb|B07869.1|B07869 [2058769] gi|2058768|gb|B07868.1|B07868~[2058768]gi|2058767|gb|B07867.1|B07867 [2058767] gi|2058766|gb|B07866.1|B07866 [2058766] gi|2058765|gb|B07865.1|B07865 [2058765] gi|2058764|gb|B07864.1|B07864 [2058764] gi|2058763|gb|B07883.1|B07883 [2058763]

gi|2058762|gb|B07882.1|B07882 [2058762] gi|2058761|gb|B07881.1|B07881 [2058761] gi|2058760|gb|B07880.1|B07880 [2058760] gi|2058759|gb|B07879.1|B07879 [2058759] gi|2058758|gb|B07878.1|B07878 [2058758] gi|2058757|gb|B07877.1|B07877 [2058757] gi|2058756|gb|B07876.1|B07876 [2058756] gi|2058755|gb|B07875.1|B07875 [2058755] gi|2058754|gb|B07874.1|B07874 [2058754] gi|2058753|gb|B07863.1|B07863 [2058753] gi|2058752|gb|B07862.1|B07862 [2058752] gi|2058751|gb|B07861.1|B07861 [2058751] gi|2058750|gb|B07860.1|B07860 [2058750] gi|2058749|gb|B07859.1|B07859 [2058749] gi|2058748|gb|B07858.1|B07858 [2058748] gi|2058747|gb|B07857.1|B07857 [2058747] gi|2058746|gb|B07856.1|B07856 [2058746] gi|2058745|gb|B07855.1|B07855 [2058745] gi|2058744|gb|B07854.1|B07854 [2058744] gi|2058743|gb|B07853.1|B07853 [2058743] gi|2058742|gb|B07852.1|B07852 [2058742] gi|2058741|gb|B07851.1|B07851 [2058741] gi|2058740|gb|B07850.1|B07850 [2058740] gi|2947527|gb|T25933.1|T25933 [2947527] gi|2924302|emb|X81655.1|EHERMAM [2924302] gi|2664256|emb|Y12234.1|EFAS48C [2664256] gi|2879906|dbj|D85752.1|D85752 [2879906] gi|2746216|gb|AF028836.1|AF028836 [2746216] gi|2745825|gb|AF039139.1|AF039139 [2745825] gi|2696019|dbj|AB007844.1|AB007844 [2696019] gi|48999|emb|X62280.1|EHPBP5G [48999] gi|2654477|gb|U89914.1|BFU89914 [2654477] gi|43347|emb|X68646.1|EHPSRAA [43347] gi|2613034|gb|AH005624.1|SEG\_EDDH4RR [2613034] gi|2613033|gb|AF029775.1|EDDH4RR2 [2613033] gi|2613032|gb|AF029774.1|EDDH4RR1 [2613032] gi|2613031|gb|AH005623.1|SEG\_EDDHIRR [2613031] gi|2613030|gb|AF029773.1|EDDHIRR2 [2613030]

gi|2613029|gb|AF029772.1|EDDHIRR1 [2613029] gi|2613028|gb|AH005622.1|SEG\_EDH19RR [2613028] gi|2613027|gb|AF029771.1|EDH19RR2 [2613027] gi|2613026|gb|AF029770.1|EDH19RR1 [2613026] gi|2613025|gb|AH005621.1|SEG\_EDISRR [2613025] gi|2613024|gb|AF029769.1|EDISRR2 [2613024] gi|2613023|gb|AF029768.1|EDISRR1 [2613023] gi|1881226|dbj|AB001488.1|AB001488 [1881226] gi|2547160|gb|AF023104.1|AF023104 [2547160] gi|2547159|gb|AF023103.1|AF023103 [2547159] gi|2547158|gb|AF023102.1|AF023102 [2547158] gi|2547157|gb|AF023101.1|AF023101 [2547157] gi|2415383|gb|AF015775.1|AF015775 [2415383] gi|2388636|gb|U94356.1|EFU94356 [2388636] gi|2388634|gb|U94355.1|ECU94355 [2388634] gi|2340825|dbj|D26045.1|D26045 [2340825] gi|2226147|emb|Y14080.1|BSY14080 [2226147] gi|2327026|gb|U87997.1|EFU87997 [2327026] gi|2318058|gb|AF012532.1|AF012532 [2318058] gi|1848175|emb|X87189.1|EM23S5SSP [1848175] gi|1848174|emb|X87187.1|EM16S23SS [1848174] gi|1848173|emb|X87188.1|EM16S23SP [1848173] gi|1848172|emb|X87185.1|EH23S5SSP [1848172] gi|1848171|emb|X87184.1|EH16S23SS [1848171] gi|1848170|emb|X87181.1|EF23S5SSP [1848170] gi|1848169|emb|X87183.1|EF23S5SPA [1848169] gi|1848168|emb|X87191.1|EF23S5SAC [1848168] gi|1848167|emb|X87180.1|EF16S23SS [1848167] gi|1848166|emb|X87182.1|EF16S23SP [1848166] gi|1848165|emb|X87190.1|EF16S23SC [1848165] gi|1848164|emb|X87186.1|EF16S23SA [1848164] gi|1848156|emb|X87179.1|ED23S5SSP [1848156] gi|1848155|emb|X87178.1|ED16S23SS [1848155] gi|1848154|emb|X87177.1|ED16S23SA [1848154] gi|2274942|emb|AJ000346.1|EHNAPBC [2274942] gi|2274939|emb|AJ000042.1|EFGLS24B [2274939] gi|414575|gb|L12710.1|ENEAAC [414575] gi|2245603|gb|AF006008.1|AF006008 [2245603]

gi|2231992|gb|U94530.1|EFU94530 [2231992] gi|2231990|gb|U94529.1|EFU94529 [2231990] gi|2231988|gb|U94528.1|EFU94528 [2231988] gi|2231986|gb|U94527.1|EFU94527 [2231986] gi|2231984|gb|U94526.1|EFU94526 [2231984] gi|2231982|gb|U94525.1|ECU94525 [2231982] gi|2231980|gb|U94524.1|ECU94524 [2231980] gi|2231978|gb|U94523.1|ECU94523 [2231978] gi|2231976|gb|U94522.1|ECU94522 [2231976] gi|2231974|gb|U94521.1|ECU94521 [2231974] gi|2196685|gb|U25090.1|EFU25090 [2196685] gi|2197120|gb|AF003922.1|AF003922 [2197120] gi|2196683|gb|U25095.1|EFU25095 [2196683] gi|2196681|gb|U25094.1|EFU25094 [2196681] gi|2196679|gb|U25093.1|EFU25093 [2196679] gi|2196677|gb|U25092.1|EFU25092 [2196677] gi|2196675|gb|U25091.1|EFU25091 [2196675] gi|2196673|gb|U24682.1|EFU24682 [2196673] gi|532533|gb|U09422.1|EFU09422 [532533] gi|487271|dbj|D17462.1|ENENTP [487271] gi|468459|dbj|D28859.1|ENEPPD1 [468459] gi|440135|dbj|D16334.1|ENEATPK [440135] gi|391680|dbj|D13816.1|ENENAABS [391680] gi|1402524|dbj|D78257.1|D78257 [1402524] gi|709995|dbj|D30808.1|BACYCB20 [709995] gi|2109265|gb|U91527.1|EFU91527 [2109265] gi|1041112|dbj|D78016.1|ENEPPD1A [1041112] gi|1339880|dbj|D85392.1|ENERPA [1339880] gi|1339878|dbj|D85393.1|ENEGE1E [1339878] gi|662918|emb|Z46807.1|EHCOPAYZ [662918] gi|769796|emb|X86176.1|EFRPODDNE [769796] gi|1854638|gb|U51479.1|EGU51479 [1854638] gi|1857221|gb|U72706.1|EFU72706 [1857221] gi|1857219|gb|U72704.1|EFU72704 [1857219] gi|1857217|gb|U72705.1|ECU72705 [1857217] gi|1272655|emb|X96978.1|EFPPD1GNS [1272655] gi|1272652|emb|X96976.1|EFPLSEP1G [1272652] gi|1279406|emb|X96977.1|EFPAD1ORF [1279406] gi|1070149|emb|X93211.1|EFTNFO1 [1070149]

gi|1065723|emb|X92947.1|EFTETMGN [1065723] gi|1019639|gb|L38972.1|PH4COINJN [1019639] gi|1151151|gb|U43087.1|EFU43087 [1151151] gi|1098507|gb|U17283.1|BMU17283 [1098507] gi|1498072|gb|U64887.1|EFU64887 [1498072] gi|1498071|gb|U64886.1|EFU64886 [1498071] gi|1469783|gb|U58049.1|EHU58049 [1469783] gi|1763666|gb|U81452.1|EFU81452 [1763666] gi|624694|gb|L38973.1|PH4SEQ [624694] gi|1730458|emb|Z83305.1|EFVANRES [1730458] gi|1419498|emb|X84796.1|ECPFW4 [1419498] gi|1419497|emb|X84795.1|ECPFW3 [1419497] gi|1419496|emb|X84794.1|ECPFW1 [1419496] gi|254400|gb|S43266.1|S43266 [254400] gi|239025|gb|S66277.1|S66277 [239025] gi|1054931|gb|U38590.1|EFU38590 [1054931] gi|1244573|gb|U39788.1|EHU39788 [1244573] gi|1244571|gb|U39789.1|EGU39789 [1244571] gi|1244569|gb|U39790.1|EFU39790 [1244569] gi|1255020|gb|U39777.1|ESU39777 [1255020] gi|1255018|gb|U39775.1|EPU39775 [1255018] gi|1255016|gb|U39778.1|EDU39778 [1255016] gi|1255014|gb|U39776.1|ECU39776 [1255014] gi|1255012|gb|U39774.1|EAU39774 [1255012] gi|1619922|gb|U69267.1|IVU69267 [1619922] gi|790436|emb|X84861.1|EFEFMPBP5 [790436] gi|790434|emb|X84858.1|EFD63RPSR [790434] gi|790432|emb|X84862.1|EF721PBP5 [790432] gi|790430|emb|X84860.1|EF63RPBP5 [790430] gi|790428|emb|X84859.1|EF366PBP5 [790428] gi|1572800|gb|U70854.1|CELF38A5 [1572800] gi|1041816|gb|U17153.1|EFU17153 [1041816] gi|1086523|gb|U39859.1|EFU39859 [1086523] gi|403564|gb|U01917.1|EFU01917 [403564] gi|1515474|gb|U66286.1|EFU66286 [1515474] gi|1513068|gb|U15554.1|LMU15554 [1513068] gi|1296520|emb|X94181.1|EFENTAORF [1296520] gi|1488069|gb|U63997.1|EFU63997 [1488069] gi|1209525|gb|U35369.1|EFU35369 [1209525]

gi|1469341|gb|U30931.1|ESU30931 [1469341] gi|488331|gb|M77276.1|SYNGIP2122 [488331] gi|1046177|gb|U39733.1| [1046177] gi|1236613|gb|U49939.1|CVU49939 [1236613] gi|47491|emb|X55766.1|SS16SR5G [47491] gi|47490|emb|X55767.1|SS16SR3G [47490] gi|47061|emb|X56353.1|SFTET916 [47061] gi|49022|emb|X62755.1|SFNPRG [49022] gi|47047|emb|X17214.1|SFPASA1 [47047] gi|47044|emb|X68847.1|SFNOXAA [47044] gi|47033|emb|V01547.1|SFKANR [47033] gi|47018|emb|X02027.1|SF5SRNA [47018] gi|511044|emb|X75752.1|MP16SRNA0 [511044] gi|511043|emb|X75751.1|MP16SR243 [511043] gi|886481|emb|X82819.1|ESPLPAM [886481] gi|517387|emb|X76177.1|ES16SRR [517387] gi|472916|emb|X76913.1|EHNTPOP [472916] gi|43351|emb|X55133.1|ES16SRRN [43351] gi|1143442|emb|X92687.1|EFPBP5G [1143442] gi|963032|emb|Z50854.1|EHARPQTOU [963032] gi|886479|emb|X84818.1|EHDNAPSR [886479] gi|551437|emb|X81654.1|EHIS1216 [551437] gi|467805|emb|X78425.1|EFPBP5 [467805] gi|296721|emb|X55961.1|EFPD78 [296721] gi|287946|emb|Z19137.1|EFPTSHGN [287946] gi|49042|emb|X63285.1|EHNAKA [49042] gi|49019|emb|X62658.1|EFSEA1 [49019] gi|43337|emb|Z12296.1|EFSPREG [43337] gi|43335|emb|X56895.1|EFPVANAG [43335] gi|43333|emb|X16421.1|EFPF54 [43333] gi|43331|emb|X62657.1|EFORF3 [43331] gi|1065721|emb|X92945.1|EFCAT501 [1065721] gi|806551|emb|Z49243.1|EF4110SOD [806551] gi|806549|emb|Z49244.1|EF4105SOD [806549] gi|505530|emb|X79542.1|EFAS48 [505530] gi|43323|emb|X62656.1|EFASP1-[43323]\_ gi|40840|emb|X56422.1|EC16SRNAG [40840] gi|48189|emb|X04388.1|TN1545TR [48189] gi|928814|gb|L40841.1|ENETRANSPO [928814] gi|141856|gb|L01794.1|AD1REPABC [141856]

gi|149125|gb|M90647.1|IP8VANY [149125] gi|141862|gb|M87836.1|AD1TRAE1 [141862] gi|141860|gb|M84374.1|AD1TRAA [141860] gi|141853|gb|M62888.1|AD1PAD1 [141853] gi|1101637|dbj|D31674.1|EVM16RNA7 [1101637] gi|1101636|dbj|D31675.1|ENE16RNA8 [1101636] gi|497792|dbj|D31676.1|ENC16RNA9 [497792] gi|1022729|gb|U36195.1|EFU36195 [1022729] gi|488338|gb|M77279.1|SYNGIP3124 [488338] gi|488335|gb|M77278.1|SYNGIP2563 [488335] gi|488333|gb|M77277.1|SYNGIP2124 [488333] gi|488329|gb|M77275.1|SYNGIP2121 [488329] gi|388267|gb|L19532.1|AD1TRAC [388267] gi|493016|gb|U03756.1|EFU03756 [493016] gi|453536|gb|L28754.1|INSTRAN [453536] gi|153658|gb|M58002.1|STRHYDROLA [153658] gi|475427|gb|U00681.1|EFU00681 [475427] gi|818704|gb|U24692.1|EFU24692 [818704] gi|155036|gb|M97297.1|TRNVAN [155036] gi|150552|gb|M64978.1|PCFPRGAB [150552] gi|786274|gb|U22541.1|EHU22541 [786274] gi|786273|gb|U22540.1|EHU22540 [786273] gi|559858|gb|L37110.1|AD1CLYL [559858] gi|643614|gb|U16659.1|ECU16659 [643614] gi|643612|gb|U16658.1|ECU16658 [643612] gi|290641|gb|L13292.1|ENECOPPUMP [290641] gi|624701|gb|L29639.1|ENEVANCRF [624701] gi|624699|gb|L29638.1|ENEVANCR [624699] gi|624692|gb|L29641.1|ENEDDLA [624692] gi|624690|gb|L29640.1|ENEDDL [624690] gi|493094|gb|L32813.1|ENERRD [493094]

[153852] gi|153851|gb|M22645.1|STRTN9162 [153851] gi|153850|gb|M20864.1|STRTN9161 [153850] gi|153660|gb|M36878.1|STRIF2BA [153660] gi|153585|gb|M13771.1|STRBRP [153585] gi|153575|gb|M64265.1|STRATPEFHA [153575] gi|153565|gb|M90060.1|STRATPASEA [153565] gi|152969|gb|M92376.1|STABLAIA [152969] gi|309660|gb|L14285.1|PCFPRGWZY [309660] gi|433714|gb|L12033.1|ENESATA [433714] gi|290645|gb|L15304.1|ENEVANB2A [290645] gi[148331|gb|M84146.1|ENEVANR [148331] gi|148329|gb|M64304.1|ENEVANH [148329] gi|148326|gb|M68910.1|ENEVANCRES [148326] gi|148324|gb|M75132.1|ENEVANC [148324] gi|148323|gb|L06138.1|ENEVANB [148323] gi|148321|gb|M85225.1|ENETETM [148321] gi|148320|gb|L00925.1|ENERTRNA [148320] gi|148319|gb|L00924.1|ENERRNA [148319] gi|148317|gb|M81466.1|ENERECA [148317] gi|148315|gb|M81961.1|ENENAPA [148315] gi|148312|gb|M38386.1|ENEMSPDPS [148312] gi|148310|gb|M37185.1|ENEGELE [148310] gi|148307|gb|L07892.1|ENEBLACREG [148307] gi|148305|gb|M60253.1|ENEBELAA [148305] gi|148303|gb|M77639.1|ENEB14NAM [148303] gi|290644|gb|L16515.1|ENERGTG [290644] gi|154954|gb|M37184.1|TRN916 [154954] gi|148301|gb|M69221.1|ENEAAD9A [148301] gi|148308|gb|M38052.1|ENECYLB [148308]

gi|153852|gb|AH000939.1|SEG STRTN916

Table 28

# Phage Dp1 complete genome sequence. 56506 nucleotides.

1	ataataaaaa	tatgaagcag	atattgggtt	aattattgct	taacaaaatg	caccgaattt	gtgtataata
71	taagtgaagc	agttttgtaa	acctgacatc	ctgctaaata	aaaataaagg	aggctcgaac	atgagtcaaa
141	acactacacg	cactgacgct	gaattgacag	gcgttactct	tttaggaaac	caagacacca	aatacgatta
211	tgactataat	ccagacgtcc	ttgaaacttt	ccctaacaaa	catcctgaaa	ataattacct	agtaacattt
281	gacggatatg	aattcacttc	cctttgccct	aaaacaggac	agcctgactt	cgcgaatgtt	ttcattagtt
351	acattccaaa	cgaaaagatg	gttgaatcta	aatcattgaa	attgtactta	ttcagtttcc	gtaaccacgg
421	tgacttccac	gaagattgca	tgaacattat	tttgaatgac	ttgtatgaat	tgatggaacc	taagtacatt
491	gaagtcatgg	gcctattcac	tectegtggt	ggaatttcaa	tttacccatt	cgtcaacaaa	gtgaatcctc
561	aatttgcaac	tcctgaactt	gaacagcttc	aacttcaacg	caaattgaac	ttccttggaa	atgttcaagg
631	tcttggacga	gctattcgat	aggaggctgg	aatgaaatca	gtagttttat	tatccggcgg	agtcgactca
701	gccacttgtt	tagcaattga	agttgacaag	tggggttcta	aaaatgttca	tgctatagca	ttcaattacg
771	gacaaaagca	tgaagcagaa	cttgaaaatg	ctgctaatgt	tgcaatgttc	tacggagtca	agttcaccat
841	tcttgaaatt	gactcgaaaa	tctactcaag	ctctagctct	teettattae	aaggaaaagg	cgaaatttca
911	catggaaaat	cttacgctga	aatcctagca	gagaaggaag	tagttgacac	ctatgttcca	tttagaaatg
981	gactaatgct	ttcacaggct	gcggcttatg	cttattcggt	tggagettet	tacgtcgtat	atggtgctca
1051	cgcagacgat	gcggctggag	gtgcttaccc	tgattgcact	cctgagttct	ataattcaat	gtcaaatgca
1121	atggaatatg	gaactggagg	caaggtaacc	cttgtcgctc	ctctacttac	tctaaccaag	gcgcaagtcg
1191	ttaaatgggg	aattgattta	gatgttcctt	atttettgae	tegtteatgt	tatgaaagtg	acgctgaaag
1261	ttgtggaact	tgcgcaactt	gracegaceg	caaaaaggca	ttegaagaaa	atggaatgac	tgaccctatt
1331						cageteatea	
1401						ttcattagca	
1471	atgaccacgg	ttcgagtcaa	gggatggttg	aggantan	ccacgccaag	aaaatcgcag	gracattear
1541						tagcaaatgc	
1611						ccttacctgg	
1681						cctacaggtt taacctttat	
1751	tacttactac	gagacteca	aattttagag	cannancana	ataatootta	atcaatacaa	tragerteaa
1821	gaaaagatta	tteesatesa	tatteacase	cctgagaaaa	tocctatcat	ggaaattttc	ggtcctacaa
1891 1961	ttesacetes	acceptacta	atacotcasa	agactatttt	cattogaact	ggtggatgcg	actateatte
2031	cccaaggiga	aggaatggtt	tracctooaa	cogtactact	gagccggaat	atatcacagg	caaagaaget
2101						taaccacgtg	
2171	geragregaa	toccttaate	aacgagccta	tooctaagat	gatttcgatt	ctaaaagaac	atggattcaa
2241	gaggaaaccc	gaaactcaag	gaactcgatt	ccaagaatgg	ttcaaagaag	taagcgatat	cactattagt
2311	cotasaccoc	crrcaagtgg	aatgagaact	aatatgaaaa	ttcttgaage	tattgtagat	agaatgaatg
2381	atgaaaacct	tgactggtca	tttaaaatcq	ttatctttqa	cqaaaatqac	ctagcttatg	cgcgtgatat
2451						ggaatgcaaa	
2521						taaagtgtat	
2591						aataaaagag	
2661						ggatgggctt	
2731	ggatgaaatt	gtaaccttgg	acaatactga	ggcagccgtt	caaagacttt	ttggtctatt	aggcgaggac
2801	gcagaacgtg	acgggttgca	agatactcca	ttccgttttg	ttaaagcact	cgctgaacat	accgtagggt
2871						gaagaccttg	
2941	agacattcca	ttcaattctt	tatgtgagca	tcatttagct	ccgttcgtag	ggaaggtgca	tattgcatac
3011	attectaagg	ataagattac	aggtctttca	aaattcggtc	gagtggttga	aggatacgct	aaacgacttc
3081	aagtacaaga	gcgcttgact	caacaaatcg	ctgacgctat	tcaggaagtt	ctaaatcctc	aagcagttgc
3151						agcacggggc	
3221	acttcaacta	tgcgaggtct	tttccaagat	gacgcatctg	ctcgagcaga	attgcttcag	ttgattaaaa
3291	agtaggaggc	ggaaaatgaa	taaaagtgca	accttttggc	ttgttcgaac	agctcttatt	gcggctctat
3361	atgtgacatt	gaccgttgca	ttttctgcta	ttagttatgg	acctattcaa	tttagagtca	gtgaagcctt
3431	gattcttcta	cctttatgga	accatagatg	gactccgggg	attgtattag	gaacaattat	tgcaaacttc
3501	ttttcacctc	ttggactgat	tgacgtttta	ttcggttcac	ttgctacctt	ccttggagta	gtggcaatgg
3571	tgaaagttgc	taagatggca	agtcctctat	attcacttat	ctgtccagtt	cttgctaatg	cttaccttat
3641	tgcgctggaa	cttcgaatag	tttactcttt	acctttttgg	gaatctgtca	tctatgtagg	aattagtgaa
3711	gcgattatcg	ttttaatttc	atacttcctt	atttccacgc	tggcgaagaa	caatcattt	agaacactga
3781	taggagcgaa	aaatgggatt	taatctatac	ttcgcaggag	gtcacgctat	tagcactgac	gattatttga
3851	aggaaagagg	agccaatcgc	ctattcaatc	aactgtacga	aagaaacggg	attggcaaaa	ggtggattga
3921	gcataagaaa	accaatccaa	gcactacttc	aaaactattc	gtcgactcta	gtgcatattc	tgctcatacc
3991	aaaggggctg	aagttgacat	tgacgcctat	accgaatacg	tgaatgataa	cgtgggaatg	tttgactgta
4061	tcgccgaact	cgataaaatt	cctggtgtat	ccagacagec	caagacacgt	gaacagcttt	cggaagcacc
4131						aagacaagct	
4201	ttccatatgg	gagaagactt	taaatggctc	aacttgatge	ccgaaactac	attcgaaggc	ggaaagcata
4271	ttccttacat	tggaatttca	ccagccaatg	actogactac	gaagcataaa	gacaagtgga	Lygaaagagt
4341	attcgaagtt	actegaaaca	gccccaaccc	ayacyccaag	actuacycat	ttgggatgac	agetactage
4411	caattagagc	gccacccatt	ccacagegee	gactetactt	togaccast	cacaggagcg	atgggaaaca
4481	ttatgacgtc	aaaaggatta	gctgacttgt	tategaagaa	ectomecocc	gatgctgtcc attttagcct	graggergee
4551	aaaaccggct	caagetgada	aggattatta	aatottcast	acatoctoss	ttgggcagag	agagudatud
4621	gccgaggact	tagasteet	caacatcasc	tatttragar	aagagetete	cgctcttatt	ttttttaaaa
4691	ccaagggaac	Ladadacty	Laucyttydt	ugat	~~~~~	-g-coccart	·····caaad

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aaaaatgaac tttttataca aaaacgcttg actttattca ctcattatcg tataatcata atataaataa 4761 4831 aacgaataag aggtaaataa aatgacagca gttcaacaag ttaagttcta cttagaagaa gccggcgctc actttctaaa agatgttgag tacagtgaca acttagagca agcaattatg aaagatattc ttaaatggaa 4901 tggcgctcat agagatgagc acgatatgaa aataacttca tacgaagtat tatagagagg ggtaaggcta 4971 tgaaaaaagt tcaaacttat caagaatatc taaaactagt tgagttcaaa cgtcaacttt ctttaaatct 5041 tcgagaagga aaaataggag tcgatgaagc ggttattcaa ttattcacct tctatagttt caacaatatc 5111 5181 gaggaacttc ctttcattgt actcaaaatg caagaggctg ccgtgaacgg gacttatgaa gcaaaactca 5251 atatgettaa aagatttaaa attatttaga aaeggettta caaaetegeg ataattegtg tatattatat atatcaaaaa aaggaggete atattatgag tattaagtte aaaaccgaag aactttcaaa aattgtttet 5321 cageteaata agttgaagee tageaagttg etagaaatea caaactattg geatattttt ggtgaeggeg aatgegteat gtttacageg tatgatgget caaactteet tegatgeatt ategaeageg atgttgaaat 5391 5461 tgacgtgatt gtgaaagcag agcagtttgg aaaacttgta gaaaagacca cggccgcaac cgtcacatta 5531 gttcctgaag 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cagttaaaat ggcagaaaag atttccagct tgcccaatgt agtcgagacg 45361 tottotaata acttogaact accttataag tatttoaata atgttataga ogototagat gaatgggago 45431 ttcacatctt cggcgaactt gataaagatg ttcaagacta cattgactct cgaaaccgaa tagcttcttc 45501 aagcaatgag cagttttegt teaagactac tecattegeg caccaggttg aatgtttega atacgcacaa 45571 gagcatccat gtttcctttt aggcgatgag caaggtttag ggaaaactaa acaggcaatt gatattgcag ttagcaggaa ggcaagtttc aaacattgtt taatcgtatg ttgcatatca gggctcaaat ggaattgggc 45641 45711 aaaagaagta ggtattcatt caaatgagtc agctcatatt ttaggaagtc gagtcactaa agatgggaaa 45781 tragtgattg acggagtttc taaacgggca gaagacttgc ttggtggcca cgacgaattc ttccttatca 45851 45921 ctaacattga aactcttcgc gatgctgtgt tcattaaata cttaaatgaa ctgacaaaaa gcggagaaat 45991 aagetecaaa gttattacaa gatgggaett acaggaacte etetaatgaa taacecaate gatgtattea 46061 atgttatgaa gtggctaggg gcggaacatc atacactgac tcagttcaaa gagcgatact gtatcgtcga 46131 ccagttcaat caaatcactg gatatcgaaa tctagctgaa cttcgcgagc ttgtcaacga ctacatgctt 46201 agaagaacga 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gaaaggacgg caaccagaat tttaaaactc gacaaactgc tcaacaaaga gcaatgctca 47111 ataatagaaa ggtatataaa tgaaattcac tgaaggaaaa aattggtata aagttggaga gatatgtcaa 47181 atgttgaacc gctctctatc tacgattaat gtttggtatg aagcaaaaga cttcgctgaa gaaaataaca 47251

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aatgtttgta gtgtcagctc aactattgac tgagttcggc gactataatt attttcaaac 49001 49071 catgcaagaa tttctcgaac gtttcgagcg ccttaagact tgtgagctat tagtcataga cgaaataggt ggaggttect taaccaagge ctettateet tatetgtatg acttggttaa ttatagggtt gacaataact 49141 tgtcgactat ttatacgact aattatactg acgatgaaat tattgacctt ttaggccaaa ggctttatag 49211 togtatatat gatacttcag tggttctaga ttttcaggca agcaatgtaa gaggattgga ggtaagcgaa 49281 attgaatcat agatatagta acatcacaac tattttctt tggcagattg tctttctttg tatttgctgc 49351 49421 geggtgteet attgtgeagg agtgeataat gagegagagt eteaagataa ggtgatteaa agttataage agaaagaaaa gtcagccgtc tacttgacag tcgatagttc aggagcttgg ctaggaagtg ctccgggagc 49491 caaggaaagt cctctctaca atgaaaaggg acagcatgta ggaaaattga aagaggtggg agagtgatac 49561 agetteaagt ettaaataaa gttetegaag aaaagagett ateeattta gaaaataatg gaattgacea 49631 agaatacttc acggattatt tagacgagta tcaatttatt caagaacact tttcgagata tggaagagtt 49701 49771 ccggacgacg aaactattct cgaccatttt cctggattcg aatttttcga aattggcgaa actgatgaat accttatega 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tcaaaatgct agcagagtta tcgctatgaa gcgtgacgaa aaatccggca tacttgaact atctgtcgtt aaaaaccgat atggcgaaga ccgaaaaatc atcgaatata tgtgggacgt tgaaactgga 50611 50681 50751 50821 acctatactc ttataggatt caaagaggaa ggcgaagaag gaactgaaaa aggcgaaagc tctccattga aagcaaaagc ctctaggtcg actgctcgtc ttcgaagtaa ggttacaagg gaaggagttg aagcattttg 50891 atgaaagtaa atggtettea aattgaageg acteetgaac aaataattga aaaacttteg agacaacttg 50961 aagacgaagg aacattcatt tttagacgaa ctaagtcgct tggaagcaac tatcaattct catgcccgtt 51031 tcatgcagga gggactgaaa agcatccctc ttgtggcatg agtagaaatc cttcttattc aggaagtaag 51101 gtgacggaag ctggaacggt tcactgtttc acttgcggct acacttcagg actaactgaa ttcgtctcga 51171 51241 atgtattagg tcgaaacgat ggagggttct atggaaacca gtggctgaaa aggaattttg gaacatctag 51311 cgaagtagtt aggcaaggcg tcagccctga agcgtttcga agaaatggga gaactgaaaa agtcgagcat 51381 aaaatcattc ctgaagagga acttgataaa taccggttta ttcatcctta tatgtatgaa cggaaattga cggacgaget categagatg tttgatgtag gttatgacaa actgcatgat tgcatcacet ttccagtacg 51451 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Table 29

### Phage dp1 ORFs list

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nb	Name	Frame	Position	Size (a.a.)	Key words
1	dp1ORF001	2	3669840390	1230	Putative tail;
2	dp1ORF002	1	3238635835	1149	Tail;
3	dp1ORF003	3	5353855877	779	DNA polymerase I;
4	dp1ORF004	3	4040142440	679	Minor structural;
5	dp1ORF005	1	2367425434	586	CM//CME Helicocci
6	dp1ORF006	2	4529646987 2223023621	563 463	SWI/SNF Helicase; Terminase;
7	dp1ORF007	3	4962450961	445	DNAb Helicase;
8	dp1ORF008 dp1ORF009	2	1316014404	414	Dividi Helidase,
10	dp1ORF010	2	86999859	386	RecA;
11	dp1ORF011	3	2801729096	359	Major head;
12	dp1ORF012	3	53466419	357	DNA pol. III beta;
13	dp1ORF013	3	1021511240	341	DNA pol. Ill gamma and tau;
14	dp1ORF014	3	5096151974	337	DNA primase;
15	dp1ORF015	1	37934728	311	
16	dp1ORF016	3	4341344303	296	Amidase;
17	dp1ORF017	1	1124212081	279	
18	dp1ORF018	3	3584736686	279	
19	dp1ORF019	2	1216112967	268	
20	dp1ORF020	1	18642658	264	exsD; Coenzyme PQQ;
21	dp1ORF021	2	25043295	263	GTP cyclohydrolase;
22	dp1ORF022	2	3089631675	259	
23	dp1ORF023	2	64197195	258 250	
24	dp10RF025	-1 3	1802618778 2599226738	248	
25	dp1ORF024	2	2151222252	246	
26 27	dp1ORF026 dp1ORF027	1	5276253490	242	
28	dp1ORF028	3	4459545299	234	
29	dp10RF029	2	6621348	228	exsB;
30	dp1ORF031	3	2694327611	222	
31	dp1ORF030	-2	1942320088	221	
32	dp1ORF032	1	5203352647	204	
33	dp1ORF033	2	76708239	189	
34	dp1ORF035	-1	1685917425	188	
35	dp1ORF036	1	4880849362	184	DNAc replication;
36	dp1ORF037	1	5585556388	177	
37	dp1ORF034	2	131652	173	
38	dp1ORF038	3	13501871	173	exsC; 6-pyruvoyltetrahydropterin;
39	dp1ORF039	3	33063803	165	Citrulline biosynthesis;
40	dp1ORF040	1 1	71927683	163	dUTPase:
41	dp10RF041	<u>3</u>	82088699 4808248561	163 159	dorrase,
42	dp1ORF042		3169932154	151	
44	dp1ORF043 dp1ORF044	-1	2521125666	151	
45	dp1ORF045	2	2534025777	145	
46	dp1ORF046	$\frac{2}{3}$	4277443202	142	
47	dp1ORF047	1	4754247961	139	
48	dp1ORF048	-3	1630816709	133	
49	dp1ORF049	-3	4362044018	132	
50	dp1ORF050	3	1508115476	131	
51	dp1ORF051	2	2976530154	129	
52	dp1ORF053	-3	4991750300	127	
53	dp1ORF052	3	3051630893	125	
54	dp1ORF054	2	1442314800	125	
55	dp1ORF055	3	2762728004	125	
56	dp1ORF056	-3	1878019151	123	
57	dp1ORF057	1	985910218	119	*
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59	dp1ORF059	-2	3771738070	117	
60 61	dp1ORF062	-2	4494045284	114	
62	dp1ORF063	1	4720047541	113	
63	dp1ORF064	2	2910829449	113	
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65	dp1ORF067	-1	4473545061	108	
66	dp1ORF068	3	2945129768	105	
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70	dp1ORF071	3	3890439209	101	
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76	dp1ORF077	1	1480015084	94	
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83	dp10RF065	-3	5124651497	83	
84	dp10RF085	-3	1060210847	81	
85	dp10RF087	-2	2979430036	80	
86	dp10RF088	3	50405279	79	
87	dp1ORF089	-2	1225612495	79	
88	dp10RF273	3	5625656486	76	
89	dp10RF078	-3	1728017507	75	
90	dp1ORF090	1	2703727261	74	
91	dp10RF091	1	4318943413	74	Holin:
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93	dp1ORF093	-2	4553845756	72	
94	dp1ORF095	3	88779089	70	
95	dp1ORF096	-1	4646946681	70	
96	dp10RF097	-1	3888839100	70	
97	dp1ORF098	1	4362743836	69	
98	dp10RF099	3	3829838507	69	
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100	dp10RF101	2	1922019426	68	
101	dp1ORF094	1	82818484	67	
102	dp10RF102	2	40344237	67	
103	dp10RF104	-1	2122421427	67	
104	dp10RF105	-2	18282028	66	
105	dp10RF106	-3	1032910529	66	
106	dp10RF108	-1	4925049447	65	
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162	dp10RF163	3	4022440367	47	
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203	dp1ORF202	2	4448344608	41	
204	dp10RF203	-3	2265622781	41	
205	dp10RF204	1	14711593	40	
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237         dp10RF236         -1         37418.37528         36           238         dp10RF237         -1         1568.1678         36           239         dp10RF238         -3         1191.1301         36           240         dp10RF239         1         26521.26628         35           241         dp10RF240         1         41893.42000         35           242         dp10RF241         -1         46913.47020         35           243         dp10RF242         -1         41231.41338         35           244         dp10RF243         -2         51199.51306         35           245         dp10RF244         -3         26976.27083         35           246         dp10RF245         -3         6171.6278         35           247         dp10RF246         -3         2724.2831         35           248         dp10RF247         1         29641.29745         34           249         dp10RF248         1         53560.53664         34           250         dp10RF250         2         23837.23941         34           251         dp10RF251         -1         39101.39205         34
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246         dp10RF245         -3         6171.6278         35           247         dp10RF246         -3         2724.2831         35           248         dp10RF247         1         29641.29745         34           249         dp10RF248         1         53560.53664         34           250         dp10RF249         2         2012.2116         34           251         dp10RF250         2         23837.23941         34           252         dp10RF251         -1         39101.39205         34           253         dp10RF252         -2         54667.54771         34           253         dp10RF253         -3         56151.56255         34           255         dp10RF254         -3         48375.48479         34           256         dp10RF255         -3         9468.9572         34           257         dp10RF256         1         15289.15390         33
246         dp10RF245         -3         61716278         35           247         dp10RF246         -3         27242831         35           248         dp10RF247         1         2964129745         34           249         dp10RF248         1         5356053664         34           250         dp10RF249         2         20122116         34           251         dp10RF250         2         2383723941         34           252         dp10RF251         -1         3910139205         34           253         dp10RF252         -2         2466754771         34           253         dp10RF253         -3         5615156255         34           255         dp10RF254         -3         4837548479         34           256         dp10RF255         -3         94689572         34           257         dp10RF256         1         1528915390         33
247         dp10RF246         -3         2724.2831         35           248         dp10RF247         1         29641.29745         34           249         dp10RF248         1         53560.53664         34           250         dp10RF249         2         2012.2116         34           251         dp10RF250         2         23837.23941         34           252         dp10RF251         -1         39101.39205         34           253         dp10RF252         -2         54667.54771         34           254         dp10RF253         -3         56151.56255         34           255         dp10RF254         -3         48375.48479         34           256         dp10RF255         -3         9468.9572         34           257         dp10RF256         1         15289.15390         33
248         dp10RF247         1         2964129745         34           249         dp10RF248         1         5356053664         34           250         dp10RF249         2         20122116         34           251         dp10RF250         2         2383723941         34           252         dp10RF251         -1         3910139205         34           253         dp10RF252         -2         5466754771         34           254         dp10RF253         -3         5615156255         34           255         dp10RF254         -3         4837548479         34           256         dp10RF255         -3         94689572         34           257         dp10RF256         1         1528915390         33
249         dp10RF248         1         53560.53664         34           250         dp10RF249         2         2012.2116         34           251         dp10RF250         2         23837.23941         34           252         dp10RF251         -1         39101.39205         34           253         dp10RF252         -2         54667.54771         34           254         dp10RF253         -3         56151.56255         34           255         dp10RF254         -3         48375.48479         34           256         dp10RF255         -3         9468.9572         34           257         dp10RF256         1         15289.15390         33
250         dp10RF249         2         2012.2116         34           251         dp10RF250         2         23837.23941         34           252         dp10RF251         -1         39101.39205         34           253         dp10RF252         -2         54667.54771         34           254         dp10RF253         -3         5615156255         34           255         dp10RF254         -3         4837548479         34           256         dp10RF255         -3         94689572         34           257         dp10RF256         1         1528915390         33
251         dp10RF250         2         2383723941         34           252         dp10RF251         -1         3910139205         34           253         dp10RF252         -2         5466754771         34           254         dp10RF253         -3         5615156255         34           255         dp10RF254         -3         4837548479         34           256         dp10RF255         -3         94689572         34           257         dp10RF256         1         1528915390         33
252         dp10RF251         -1         3910139205         34           253         dp10RF252         -2         5466754771         34           254         dp10RF253         -3         5615156255         34           255         dp10RF254         -3         4837548479         34           256         dp10RF255         -3         94689572         34           257         dp10RF256         1         1528915390         33
253     dp10RF252     -2     5466754771     34       254     dp10RF253     -3     5615156255     34       255     dp10RF254     -3     4837548479     34       256     dp10RF255     -3     94689572     34       257     dp10RF256     1     1528915390     33
253     dp10RF252     -2     5466754771     34       254     dp10RF253     -3     5615156255     34       255     dp10RF254     -3     4837548479     34       256     dp10RF255     -3     94689572     34       257     dp10RF256     1     1528915390     33
254     dp1ORF253     -3     5615156255     34       255     dp1ORF254     -3     4837548479     34       256     dp1ORF255     -3     94689572     34       257     dp1ORF256     1     1528915390     33
255         dp10RF254         -3         48375.48479         34           256         dp10RF255         -3         9468.9572         34           257         dp10RF256         1         1528915390         33
256 dp10RF255 -3 9468.9572 34 257 dp10RF256 1 1528915390 33
257 dp10RF256 1 1528915390 33
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259 dp10RF258 1 4402344124 33
260 dp10RF259 2 4298.4399 33
261 dp10RF260 2 2474624847 33
262 dp10RF261 3 288.389 33
263 dp10RF262 3 94089509 33
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203   dp10KF204   41   0000::0103   00
266 dp10RF265 -1 47004801 33
267 dp10RF266 -2 5011950220 33
268 dp10RF267 -2 47266.47367 33
269 dp10RF268 -2 1252012621 33
270 dp10RF269 -3 5373353834 33
271 dp10RF270 -3 5069150792 33 -

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38882

38966

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37874

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38042

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37706

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### Table 30

### Predicted Dp-1 amino acid sequences

atgattgacaataatttacctatgagtccaattcctggcgaaattgttcaagtatatgaccaaaacttcaatctaattggagca MIDNNLPMSPIPGEIVQVYDQNFNLIGA agtgatgaaatctttagcaagcattacgaagacgaaattgtgactcgagctcgaggaaaagaaactttcacttttgaaagtatt S D E I F S K H Y E D E I V T R A R G K E T F T F E S I qaaacctcatctatctatcaacacttaaaggttgaaaacattatccagtatggaggaagatggtttcgaattaaatatgctcag ETSSIYQHLKVENIIQYGGRWFRIKYAQ gacgtagaagatgtcaaagggcttaccaagtttacctgctacgcattatggtatgaactagcagaaggcttgcctaggaagttg V E D V K G L T K F T C Y A L W Y E L A E G L P R K L aaacacgttgcttcttctgtaggcgctgtcgcgctagatattatcaaagacgcaggtgaatgggttcgactagtttgtcctcct K H V A S S V G A V A L D I I K D A G E W V R L V C qacqqtqctaacaacaagttcqaagcataacagccgcagaaaattcaatgcttttggcatcttcgatatcttgcaaagcaatac G A N K Q V R S I T A A E N S M L W H L R Y L A K Q Y gtcgagtctaaagtagactttcctcttgtagttgaagagaatttgaaatatgtcactaggcaggaagattctcgaaacctgtgt V E S K V D P P L V V E E N L K Y V T R Q E D S R N L C acggcttacaagttgacaggtaaaaaggaagaaggcagtcaagagcctttaacgtttgcttctatcaacaatggaagtgaatatAYKLTGKKEEGSQEPLTFASINNGSEY ctcattgatgtttcgtggtttactacacgccacatgaagcctcgatatattgctaaatctaaaagcgacgaacattttagaatt V S W F T T R H M K P R Y I A K S K S D E H F R I K E N L M S A A R A Y L D I Y S R P L I G Y E A S A  ${\tt tataacaaggttcctgacttgcatcatactcaactaattgtcgacgaccattatgatgttatcgagtggcgaaagatatctgct}$ YNKVPDLHHTQLIVDDHYDVIEWRKISA cgaaaaattgactacgacgacctttcaaactctactatcattttccaagaccctcgaaaagacttgatggacttgctaaatgag R K I D Y D D L S N S T I I F Q D P R K D L M D L L N E gacggcgaaggagtcctttcaggggaaactgtaaatgagtcccaagttgttattagatacgcagatgacattttagggactaatD G E G V L S G E T V N E S Q V V I R Y A D D I L G T N tttaatgcagaatctgggaaatacattggtgtccttaatactaataagaaaccgagcgaattagttcctgacgactttacatgg PNAESGKYIG VLNTNKKPSEL V PDD FT W attcgactagaaggtcctaaaggtgacgcaggtttaccgggagctcctgggcgtgatggagtcgacggtgtacctggaaagagc R L E G P K G D A G L P G A P G R D G V D G V P G K S V G I A D T A I T Y A V S V S G T Q E P E N G W S E  $\tt gttcctgaactcataaaaggtcgattcttgtggactaaaacattttggagatatactgacggctcacatgaaactggatactcc$ PELIKGRFLWTKTFWRYTDGSHETG gttgcctatatagggcaagacggaaattccggaaaagacggaatcgcaggtaaggacggagtaggtatagccgcaactgaagtc V A Y I G Q D G N S G K D G I A G K D G V G I A A T E V  ${\tt atgtatgcaagttcgccatctgctactgaagctccagctggttggattggtctacgcaagttcctaccgtcccaggttggtcagtat}$ M Y A S S P S A T E A P A G G W S T Q V P T V P G G Q Y ttatggactcgaacaagatggcgctacactgaccaaactgatgaaattggatattcagtttcaagaatgggcgagcagggtcct L W T R T R W R Y T D Q T D B I G Y S V S R M G E Q G P aaaggtgacgcaggtcgtgacggtattgcaggaaagaacggaatagggttgaagtcaacttcagtttcttatggaattagtcccKGDAGRDGIAGKNGIGLKSTSVSYGISP  ${\tt actgattctgcgattcctggagtatgggcttcacaagttccttctttaatcaaaggtcaatatctttggactcgaactattttgg$ G V W A S Q V P S L I K G Q Y L W T R T I acctataccgattcaactaccgaaacgggctatcaaaaaaacctacattccaaaagacgggaatgacggtaaaaatggaattgct TYTDSTTETGYQKTYIPKDGNDGKNG ggtaaggatggggtaggaattaagtctacgaccattacctacgcaggctcaacctcaggaacagttgcgcctacttcaaattgg K D G V G I K S T T I T Y A G S T S G T V A P T S N W acttctgctattccaaatgttcaaccgggattcttcttgtggacgaaaactgtttggaactatactgatgacactagcgaaaca T S A I P N V Q P G F F L W T K T V W N Y T D D T S E T ggttactcagtttccaagataggtgaaacaggtcctagaggagttctaaggtcttcaaggtcctcaagggcttcaaggaattcct G Y S V S K I G E T G P R G V Q G L Q G P Q G L Q G I P ggacctgcaggagctgacggacgttcgcaatatactcacctcgctttctctaatagtccaaacggtgagggatttagtcatact G P A G A D G R S Q Y T H L A F S N S P N G E G F S H T gacagcggacgagcatacgtcggtcagtatcaagatttcaatcccgtccattcaaaagaccctgcagcctatacatggacgaaa D S G R A Y V G Q Y Q D F N P V H S K D P A A Y T W T K tggaagggaatgacggagctcaagggatacccgggaagccaggcgagacggtaagactaattatttccatatagcttacgct K G N D G A Q G I P G K P G A D G K T N Y F H I A Y S A D G S R E F S L E D N N Q Q Y M G Y Y S D Y E Q A gatagcagggatcgaactaagtatcgatggtttgaccgccttgccaatgttcaagtgggaggtcgaaacgagttccttaattct D S R D R T K Y R W F D R L A N V Q V G G R N E F L N S L F E F G L K P R Y S S Y N L M D G Q D Q T Q G Q I S A actattgacgaacgtcaacggttcaaaggtgctaactctttacgacttgactcaacatggaacggtaaaccgcagaaccaaaaa TIDER ORFK GANSLRLDST WNG KP ON OK

39638

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40226 1177

40310 40390

1205 dp10RF002

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141 32890

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33730 449

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33898 505

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561 34150

589

617 34318

645

673

34402

34486 701

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L T F S L G G D T R L G T P T E W S N L E G R I S F W A aaggcctctaggaacggagtgagcttagctgcacggccgggttatcgtagtaacgtatttaccgcaaccttaaccgatcaatgg KĀSRNG VSLĀĀRPGYRSNVFTĀTLTDQW aagttctacgattttaaattctttgacaaagttaattcaaattgtaccgctgaagcaattttccatgtattcactcaaagttgt K F Y D F K F F D K V N S N C T A E A I F H V F T O S C  ${\tt tcagtgtggctcaatcatattaaaatcgaacttggtaatatctctactccttttagtgaagcagaggaagaccttaaatatcga}$ S V W L N H I K I E L G N I S T P F S E A E E D L K Y R  ${\tt attgactcaaaagccgatcaaaagctaactaaccaacagttgacggcactcacggaaaaggctcaactacatgacgcagaactg}$ I D S K A D Q K L T N Q Q L T A L T E K A Q L H D A E L aaagctaaggctacaatggagcagttaagtaacttagaaaaggcttatgaaggtagaatgaaagctaatgaagaagctatcaaa KAKAT MEQLSNLEKAY EGRMKANE BAIK aaatcggaagccgacctaatcttagcggcaagtcgaattgaagctactatccaagaacttggcgggctacgggaactgaagaag K S E A D L I L A A S R I E A T I Q E L G G L R E L K K ttcqtcqacaqttacatqaqctcttctaatgaaggtctaattatcggtaagaacgacggtagctctaccattaaggtatcaagt F V D S Y M S S S N E G L I I G K N D G S S T I K V S S caatccattcaagtcggccgatttagaacggaacaatactcgtttaatccagacatgaacgtgattcggtatgtaggataa

Q S I Q V G R F R T E Q Y S F N P D M N V I R Y V G \*

atggattttgggtcaattgcagcaaaaatgactttggatatctcaaacttcacaagtcaattaaatcttgctcaaagtcaagcg M D F G S I A A K M T L D I S N F T S Q L N L A Q S Q A caacggctcgcactagagtcttcgaagtcctttcaaattggttctgctttaacaggattagggaaaggacttacgactgcggtt Q R L A L E S S K S F Q I G S A L T G L G K G L T T A accettectettatgggatttgcagecgcetetattaaagtagggaatgaattccaageteaaatgtcccgtgttcaagetatt T L P L M G F A A A S I K V G N B F Q A Q M S R V Q A I gcaggagcgacagcggaagagcttggtagaatgaagactcaagcaatcgaccttggtgctaaaactgcttttagtgcaaaagag A G A T A E E L G R M K T Q A I D L G A K T A F S A K E  ${\tt gcggctcaaggtatggaaaatctagcttcagccggtttccaggtaaatgaaatcatggacgctatgccaggggtacttgacctg}$ A A O G M E N L A S A G F Q V N E I M D A M P G gctgccgtatctggaggagatgtggccgcgagctccgaggccatggctagttcacttcgagcctttggattagaggcaaaccag A A V S G G D V A A S S E A M A S S L R A F G L E A N Q gcgggtcacgtggctgacgtatttgctcgagcagcagctgatacgaacgcagaaactagcgacatggcagaggcgatgaaatac A G H V A D V F A R A A A D T N A E T S D M A E A M K Y  $\tt gtcgcacccgttgctcactctatgggcttgagccttgaagaaacggctgcgtctattgggattatggccgacgccggtattaaggcctgcactctattgggattattatggccgacgccggtattaaggcctgcactctattgggattattattggccgacgccggtattaaggcctgcactctattgggattattattggccgacgccgggtattaaggcctgcactctattgggattattattggccgacgccgggtattaaggcctgacgccgacgccgggtattaaggcctgacgccgacgccgggtattattggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgggtattaaggccgacgccgacgccgggtattaaggccgacgccgacgccgggtattaaggccgacgccacgcacgccacgccacgc$ V A P V A H S M G L S L B B T A A S I G I M A D A G I K ggctcgcaagccggaaccacgcttagaggcgctctctcgcgtattgccaaacctacgaaagcgatggtcaaatcaatgcaggaa G S Q A G T T L R G A L S R I A K P T K A M V K S M Q E ttaggagtttcgttctacgacgcgaacggaaacatgattccactaagagaacaaatcgctcaactgaaaacagctactgcagga L G V S F Y D A N G N M I P L R B Q I A Q L K T A T A G L T Q E E R N R H L V T L Y G Q N S L S G M L A L L D A G P E K L D K M T N A L V N S D G A A K E M A E T M Q D aaccttgctagtaaaatcgagcaaatgggaggagctttcgagtctgttgctattattgttcaacaaatccttgagcctgcactt N L A S K I E Q M G G A F E S V A I I V Q Q I L E P A L A K I V G A I T K V L E A F V N M S P I G Q K M V V I F geaggaatggttgeagecettggaecactgettetaattgeaggaatggtgatgaeaactattgteaagttaagaattgetattA G M V A A L G P L L I A G M V M T T I V K L R I A I cagtttttaggtccagcatttatgggaacgatgggaaccattgcaggagttatagcaatattctatgctctggtcgccgtgttc Q F L G P A F M G T M G T I A G V I A I F Y A L V A V F atgatageetacacaaaateggagagatttagaaaetttatcaacagtettgegeetgetattaaagetgggtttggaggageg M I A Y T K S E R F R N F I N S L A P A I K A G F G G A LEWLLPRLKELGEWLQKAGEKAKEFGQS V G S K V S K L L E Q F G I S I G Q A G G S I G Q F I G N V L E R L G G A F G K V G G V I S I A V S L V T K F G ctcgcatttctagggattacaggaccactcgggattgctattagtctgttagtttcattttttgacagcttgggctagaacaggt LAPLGITGPLGIAISLLVSFLTAWARTG gagttcaacgcagacggaattactcaagtattcgaaaacttgacaaacacaattcagtcgacggctgatttcatctctcaatac EFN A D G I T Q V F E N L T N T I Q S T A D F I S Q Y  $\verb"cttccagtctttgtcgaaaaaaggaactcaaattttagttaagattattgaaggaattgcatctgctgttcctcaagtagttgaagttgaagttgaagttgcatctgctgttcctcaagtagttgaag$ L P V F V B K G T Q I L V K I I B G I A S A V P Q V V B gtgatttcacaagtcattgaaaatattgtgatgacaatttcgacagttatgcctcaattagtcgaagcaggaattaagataCtb ISQVIENIVMTISTV MPQLVEAGI<del>K</del>IL qaaqcqcttataaatqqtcttqttcaatctcttcctactatcattcaagcagctqttcaaattatcactqctttattcaatqqt E A L I N G L V Q S L P T I I Q A A V Q I I T A L F N G cttgttcaggcacttcctacgcttattcaagcaggtcttcaaattttgtcagctctcataaacggactagttcaagcgcttccg L V Q A L P T L I Q A G L Q I L S A L I N G L V Q A L P gcaattattcaagcagctgttcaaattatcatgtcgcttgttcaagcactaattgaaaacttgcctatgataatcgaagcagcg Ă II Q A A V Q I I M S L V Q A L I E N L P M I I E À À

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34654 757 M Q I I M G L V N A L I E N I G P I L È À G I Q I L M A 34738 ttaatcgagggacttattcaagtgcttcctgaactaattacagcagcgattcaaatcattacttcactattagaagcaatcttg LIEGLIQVLPELITAAIQIITSLLEAIL 785 34822 tegaacetteeteaacttetagaageeggagttaaattgettttateacttetteaagggttgetaaatatgetteeteaacta S N L P Q L L E A G V K L L S L L Q G L L N M L P Q L 813 attgcagggctttgcaaatcatgatggcacttcttaaagcagttatcgacttcgtccctaaacttcttcaagcaggtgttcaa 34906 I A G A L Q I M M A L L K A V I D F V P K L L Q A G V Q 841 34990 cttcttaaggcattgattcaaggtattgcttcacttctcggctcacttttatcgacagctggaaacatgctttcatcattagtt 869 L L K A L I Q G I A S L L G S L L S T A G N M L S S L V 35074 agcaagattgctagctttgtgggacagatggtttcaggaggtgcgaacctgattcgaaacttcattagtggtattgggtcaatg S K I A S F V G Q M V S G G A N L I R N F I S G I G S M 897 attggttcagctgtctctaaaattggcagcatgggaacttcaattgtttctaaggttactggattcgctggacaaatggtaagc 35158 I G S A V S K I G S M G T S I V S K V T G F A G Q M V S 925 35242 gcaggggtcaaccttgttcgaggatttatcaatggtatcagttccatggtaagttctgcggtaagtgcggcggctaatatggct V N L V R G F I N G I S S M V S S A V S A A A N M A 953 aqcagtgcattaaatgccgttaagggattcttaggtattcactctccttcacgtgtcatggagcagatgggtatctatacgggt 35326 S S A L N A V K G F L G I H S P S R V M E Q M G I Y T G 981 35410 Q G P V N G I G N M I R T T R D K A K E M Å E T V T E Å 1009 ctcagcgacgtgaagatggatattcaagaaaatggagttatagaaaaggttaaatcagtttacgaaaagatggctgaccaactt 35494 L S D V K M D I Q E N G V I E K V K S V Y E K M A D Q L 1037 cctgaaactcttccagctcctgatttcgaagatgttcgtaaagcagccggttcgcctcgagtggacttgttcaatacaggaagt 35578 PETLPAPDFEDVRKAAGSPRVDLFNTGS 1065 gacaaccctaaccaacctcagtcacaatctaaaaacaatcaaggcgagcaaaccgttgtcaacattggaacaatcgtagttcga . 35662 1093 D N P N Q P Q S Q S K N N Q G E Q T V V N I G T I V V R 35746 1121 N N D D V D K L S R G L Y N R S K E T L S G F G N I V T 35830 ccgtaa 35835 1149

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             at gaat t cact t ccct t t gccctaaaa cag gacag cct gact t cgc gaat g t t t cat t a g t t a cat t ccaa a c gaaa a g a t g a cat t cact t ccaa a c gaaa a g a t g a cat t cact t ccaa a c g a a a g a cat t cact t cact t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t cac t
288
             M N S L P F A L K Q D S L T S R M F S L V T F Q T K R W
372
             ttgaatctaaatcattga 389
             LNLNH
29
dp10RF262
             {\tt atgcctattcaactccaggcggaaagatgtggaagcatgcttgtgcagttcgacttaaatttagaaaaggtgactaccttgacg}
9408
             MPIQLQAERCGSMLVQFDLNLEKVTTLT
             aaaacggtgcatcattga 9509
9492
             KTVHH *
dp10RF263
             {\tt atgaaaattttagcatcgagttctttcgaagttttcgaaataatttccttcacctgtttgatagttggttcatctagacctttt}
27052
             M K I L A S S S F E V F E I I S F T C L I V G S S R P F
             aacaagtcttctaattga 26951
26968
             NKSSN *
29
dp10RF264
             \tt gtgaatagtacaaggcggtctaatacgctcaggatttctgctgtagggatagccgcatcatcttcaaactcaattgagtcaagc
6139
             V N S T R R S N T L R I S A V G I A A S S S N S I E S S
1
             tgtgaaacgtcttcataa 6038
6055
29
             CETSS *
dp10RF265
4801
             V N K V K R F C I K S S F F F K K N K S E K L L S K I V
1
             gacgttgacgatttttaa 4700
4717
             D V D D F *
29
dp10RF266
50220
             atgcccgttcttccaagcagttgcaagcattttatcaatagtccacgacttaccttgtccaggtcgagccattatgacaatcaa
             M P.V L P S S C K H F I N S P R L T L S R S S H Y D N Q
             atcctcaccaggaagtaa 50119
50136
            ILTRK
29
dp10RF267
             {\tt atggtcaaggtctgttctaggttcaggaagaacaaacgggaagtgaatgttattttcttcagcgaagtcttttgcttcatacca}
47367
             MVKVCSRFRKNKREVNVIFFSEVFCFIP
             aacattaatcgtagatag 47266
47283
             NINRR *
dp10RF268
```

```
12621
12537
                    ttgtcaattctagagtaa 12520
                    LSILE
29
dp10RF269
                    {\tt gtgaatagtatcgagtccatcagtttctacgtcaatagaacctattccgtcttcaatcattttgtctacatactgctcgagttt}
53834
                    V N S I E S I S F Y V N R T Y S V F N H F V Y I L L E F
                    tgcttcctcagtgattaa 53733
53750
                    CFLSD *
29
dp10RF270
                    at gatttttcggtcttcgccatatcggtttttaacgacagatagttcaagtatgccggatttttcgtcacgcttcatagcgatagcgatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcaggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcgatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcggatagcgatagcgatagcggatagcggatagcgatagcgatagcggatagcgatagcgatagcggatagcggatagcgatagcggatagcgatagcggatagcggatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcgatagcatagcgatagcgatagcgatagcgatagcgatagcgatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagcatagca
50792
                    MIFRSSPYRFLTTDSSSMPDPSSRFIAI
                    actctgctagcattttga 50691
50708
29
                    TLLAF
dp10RF271
                    atgaggetgetttgetttatettegttacegtattgacegacttectactegegaacetteetacaagaatteataceteaaag
19739
                    M R L C F I F V T V L T D F L L A N L P T R I H T S K
                    gctttttgtcagccttag 19638
19655
29
                    A F C Q P *
dp10RF272
                    \tt gtggtcaagtctgtcaatgaatgtacctgcgattttcttgacgtgataaaagtcaacaaccatcccttgactcgaaccgtggtc
1556
                     V V K S V N B C T C D F L D V I K V N N H P L T R T V
                    ataagttccgcctgctaa 1455
1472
29
                    ISSAC
dp10RF273
                    {\tt atggatttcattaggactgagtcctcttggaattggaacggttgcatatatagatattccgtcagccgtactaggccaagttct}
56256
                    M D F I R T E S S W N W N G C I Y R Y S V S R T R P S S
                    agttcagtttatcttgcagtcaattgcttcgagatatttgaaaaagtagtcaggaaaattcctgattatcttgcagtcaattgc
56340
                    S S V Y L A V N C F E I F E K V V R K I P D Y L A V N C
29
                    56424
57
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#### Table 31

Query= sid|114822|lan|dplORF001 Phage dp1 ORF|36698-40390|2 (1230 letters) >qi|928828 (L44593) ORF1904; putative [Lactococcus lactis phage BK5-T] Length = 1904 Score = 427 bits (1086), Expect = e-118 Identities = 226/475 (47%), Positives = 281/475 (58%), Gaps = 45/475 (9%) Query: 395 AESGKYIGVLNTNKKPSELVPDDFTWIRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIAD 454 + P D+TW + +G+ G GA G+DGV A+ YIG GK GVGI Sbjct: 820 ADYPSYIGQYTDFIQYDSAKPSDYTWSLI---RGNDGKDGATGKDGV---AGKDGVGIKT 873 Query: 455 TAITYAVSVSGTQEPENGWSEQVPELIKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNS 514 T ITYA+S SGT +P GW+ QVP L+KG++LWTKT W YTD S ETGYSV YI +DGN+ Sbjct: 874 TVITYALSSSGTDKPNTGWTSQVPTLVKGQYLWTKTVWTYTDSSSETGYSVTYIAKDGNN 933 GKDGIAGKDGVGIAATEVMYASSPSATEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTD 574 Query: 515 G DGIAGKDGVGI T + YA S T APA GW++QVP VP GQ+LWT+T W YTD T Sbjct: 934 GNDGIAGKDGVGIKKTTITYAVGTSGTTAPASGWNSQVPNVPAGQFLWTKTVWTYTDNTS 993 Query: 575 EIGYSVSRMGEQGPKGDAGR---DGIAGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVP 630 E GYSV+ MG +G KGD G +GIAGK+G G+K+T+++Y SP + P G W++ VP Sbjct: 994 ETGYSVAMMGVKGDKGDPGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVP 1053 Query: 631 SLIKGQYLWTRTIWTYTDSTTETGYQKTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGST 690 + KG +LWTRTIWTYTD+TTETGY Y+ +GN+G +G GKDG GIK+TTITYAGST Sbjct: 1054 PVAKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGST 1113 Query: 691 SGTVAPTSNWTSAIPNVQPGPFLWTKTVWNYTDDTSETGYSVSKIGETXXXXXXXXXXXX 750 SGT P + WTS +P V G +LWTKTVW YTD+TSETGYSV+ +G Sbjct: 1114 SGTTPPNNGWTSTVPTVAEGNYLWTKTVWTYTDNTSETGYSVAMMG-----VKGDKGDP 1167 Query: 751 XXXXXXXXXXADGRS-QYTHLAFSNSPNGEGFSHTDSGRAYVGQYQDFNPVHSKDPAAYT 809 A G+ DG+ + T + + SPNG Sbjct: 1168 GNNGTNGIAGKDGKGIKATAITYQASPNGT------TAPTGTWSASVPPVAKGSFLWT 1219 Query: 810 WTKW------KGNDGAQGIPGKPGADGKTNYFHIAYASSADGS 846 GN+G G PGK G KT I YA S G+ Sbjct: 1220 RTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTT--TITYAGSTSGT 1272 Score = 396 bits (1007), Expect = e-109 Identities = 208/449 (46%), Positives = 260/449 (57%), Gaps = 42/449 (9%) Query: 421 IRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480 + + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP + Sbjct: 1155 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPV 1211 Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNSGKDGIAGKDGVGIAATEVMYASSPSA 540 KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S Sbjct: 1212 AKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSG 1271 Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDEIGYSVSRMGEQGPKGDAGR---DGI 597 T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G Sbjct: 1272 TTPPNNGWTSTVPTVAEGNYLWTKTVWTYTDNTSETGYSVAMMGVKGDKGDPGNNGTNGI 1331 Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKGQYLWTRTIWTYTDSTTETGYQ 656 AGK+G G+K+T+++Y SP + P G W++ VP + KG +LWTRTIWTYTD+TTETGY Sbjct: 1332 AGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPVAKGSFLWTRTIWTYTDNTTETGYA 1391 Query: 657 KTY1PKDGNDGKNGIAGKDGVGIKSTTITYAGSTSGTVAPTSNWTSA1PNVQPGFFLWTK 716 Y+ +GN+G +G GKDG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK \_\_\_\_\_ Sbjct: 1392 VAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1451 TVWNYTDDTSETGYSVSKIGETXXXXXXXXXXXXXXXXXXXXXXXADGRS-QYTHLAFSNS 775 Query: 717 TVW YTD+TSETGYSV+ +G DG+ + T + + S

Sbjct: 1452 TVWTYTDNTSETGYSVAMMG-----VKGDKGDPGNNGTNGIAGKDGKGIKATAITYQAS 1505

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Query: 776 PNGEGFSHTDSGRAYVGQYQDFNPVHSKDPAAYTWTKW------KGND 817
                 AG+ P+K +TTW
                                                                    CN+
Sbjct: 1506 PNGT------TAPTGTWSASVPPVAKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNN 1557
Query: 818 GAQGIPGKPGADGKTNYFHIAYASSADGS 846
           G G PGK G KT I YAS G+
Sbjct: 1558 GHDGFPGKDGTGIKTT--TITYAGSTSGT 1584
 Score = 384 bits (977), Expect = e-105
 Identities = 179/322 (55%), Positives = 222/322 (68%), Gaps = 7/322 (2%)
Query: 421 IRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIADTAITYAVSVSGTQRPENGWSEQVPEL 480
            + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
Sbjct: 1311 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPV 1367
Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNSGKDGIAGKDGVGIAATEVMYASSPSA 540
            KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1368 AKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSG 1427
Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDEIGYSVSRMGEQGPKGDAGR---DGI 597
           T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G
                                                                   +GI
Sbjct: 1428 TTPPNNGWTSTVPTVAEGNYLWTKTVWTYTDNTSETGYSVAMMGVKGDKGDPGNNGTNGI 1487
Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKGQYLWTRTIWTYTDSTTETGYQ 656
            AGK+G G+K+T+++Y SP + P G W++ VP + KG +LWTRTIWTYTD+TTETGY
Sbjct: 1488 AGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPVAKGSFLWTRTIWTYTDNTTETGYA 1547
Query: 657 KTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGSTSGTVAPTSNWTSAIPNVQPGFFLWTK 716
              Y+ +GN+G +G GKOG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK
Sbjct: 1548 VAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1607
Query: 717 TVWNYTDDTSETGYSVSKIGET 738
           TVW YTD++ ETGYSV K+G T
Sbjct: 1608 TVWAYTDNSFETGYSVGKMGNT 1629
 Score = 201 bits (507), Expect = 2e-50
 Identities = 121/297 (40%), Positives = 156/297 (51%), Gaps = 19/297 (6%)
Query: 421 IRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
Sbjct: 1467 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPV 1523
Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNSGKDGIAGKDGVGIAATEVMYASSPSA 540
            KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1524 AKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSG 1583
Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDEIGYSVSRMGEQGPKGDAGRDGIAGK 600
           T P GW++ VPTV G YLWT+T W YTD + E GYSV +MG GP AG +G GK
Sbjct: 1584 TTPPNNGWTSTVPTVAEGNYLWTKTVWAYTDNSFETGYSVGKMGNTGP---AGSNGNPGK 1640
Query: 601 NGIGLKSTSVSYGISPTDSAIPGVWASQVPSLIKG-QYLWTRTIWTYTDSTTE--TGYQK 657
                + T+ G++ S + + ++ G +Y W W +
Sbjct: 1641 VVSDTEPTTKFKGLTWKYSGVVDMPLGNGTKILAGTEYYWNGNNWALYEINAHNINGDNL 1700
Query: 658 TYIPKDGNDGK-NGIAGKDGVGIKSTTITYAGS----TSGTVAPTSNWTSAIPNVQ 708
+ DGK I G +GV + T T GS +S + T N T AI N Q
Sbjct: 1701 SVTNGTFKDGKIESIWGSNGV---NGTTTIEGSHLQIHSSDSTTNTEN-TLAIDNRQ 1753
Query= sid | 114823 | lan | dp1ORF002 Phage dp1 ORF | 32386-35835 | 1
         (1149 letters)
>dbj|BAA31888| (AB009866) orf 15 [bacteriophage phi PVL]
          Length = 694
 Score = 280 bits (709), Expect = 3e-74
Identities = 157/465 (33%), Positives = 257/465 (54%), Gaps = 28/465 (6%)
Query: 40 QIGSALTGLGKGLTTAVTLPLMGFAAASIKVGNEFQAQMSRVQAIAGATAEELGRMKTQA 99
          +IG+++ +G+ +T VT P++ A + K G EF M +V+A +GAT EE +K +A
Sbjct: 151 EIGNSMKNVGRNMTMYVTAPVVAGFAVAAKKGIEFDDSMRKVKATSGATGEEFEALKKKA 210
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++GA T FSA ++A+ + +A AG+ ++M+ + GV+DL
Sbjct: 211 REMGATTKFSASDSAEALNYMALAGWDSKQMMEGLSGVMDLAAASGEELGAVSDIVTDGL 270
Query: 160 RAFGLEANQAGHVADVFARAAADTNAETSDMAEAMKYVAPVAHSMGLSLEETAASIGIMA 219
           AFGL+A +GH+ADV A+ ++ N + + EA KYVAPVA ++G ++E+T+ +IG+M+
Sbjct: 271 TAFGLKAKDSGHLADVLAQTSSKANTDVRGLGEAFKYVAPVAGALGYTIEDTSIAIGLMS 330
Query: 220 DAGIKGSQAGTTLRGALSRIAKPTKAMVKSMQELGVSFYDANGNMIPLREQIAQLKTATA 279
          +AGIKG +AGT LR + ++ PT+AM M+ LG+S D+NG MIP+R+ + QL+
Sbjct: 331 NAGIKGEKAGTALRTMFTNLSSPTRAMGNEMERLGISITDSNGKMIPMRKLLDQLREKFK 390
Query: 280 GLTQEERNRHLVTLYGQNSLSGMLALLDAGPEKLDXMTNALVNSDGAAKEMAETMQDNLA 339
                   T++G+ ++SG LA+++A E K+T ++ +S GA+K MA+TM+ L
Sbjct: 391 HLSKDQQASSAATIFGKEAMSGALAIINASDEDYQKLTKSIDSSTGASKRMADTMESGLG 450
Query: 340 SKIEQMGGAFESVAIIVQQILEPALAKIVGAITKVLEAFVNMSPIGQKMVVIFAGMVAAL 399
           K+ + E +A+ + +EPAL IV A +KV+
                                               + Q VV F VA L
Sbjct: 451 GKLRTLRSQLEELALTIYDRIEPALKIIVSAPSKVVTWVTKLPTSIQLAVVGFGLFVAVL 510
Query: 400 GPLLLIAGM------ VMTTIVKLRIAIQFLGPAFMGTMGTIAGVIAIF------ 441
          GPL+ + G+
                        MT + LI +
                                         P
                                                IA ++ +F
Sbjct: 511 GPLVFMFGLFISVMGNAMTVLGPLLINVNKASGLFAFLRTKIASLVKLFPILGVSISSLT 570
Query: 442 -----YALVAV---FMIAYTKSERFRNFINSLAPAIKAGFGGA 476
                  ALV + F AY +SE FRN +N + F A
Sbjct: 571 LPITLIVGALVGIGIAFYQAYKRSETFRNIVNQAISGVANAFKAA 615
Query= sid | 114824 | lan | dp10RF003 Phage dp1 ORF | 53538-55877 | 3
        (779 letters)
>ap|P43741|DP01_HABIN DNA POLYMERASE I (POL I) >gi|1074025|pir||E64098 DNA polymerase I
          (polA) homolog - Haemophilus influenzae (strain Rd KW20)
          >gi|1573871 (U32767) DNA polymerase I (polA)
          [Haemophilus influenzae Rd]
          Length = 930
 Score = 191 bits (481), Expect = 1e-47
 Identities = 148/553 (26%), Positives = 262/553 (46%), Gaps = 60/553 (10%)
Query: 63 RLELITEEAKLEQYVDKMIEDGIGSIDVETDGLDTIHDELAGVCLYSPSQKGIYAPVNHV 122
+ E + +A L ++++K+ + ++D ETD LD + L G+ + + Y P+

Sbjct: 333 KYETILTQADLTRWIEKLNAAKLIAVDTETDSLDYMSANLVGISFALENGEAAYLPLQLD 392
Query: 123 SNMTKMRIKNQISPEFMKKMLQRIVDSGIPVIYHNSKFDMKSIYWRLGVKMNEPAWDTYL 182
++ + +K +L+ + I I N KFD +SI+ R G+++ +DT L
Sbjct: 393 YLDAPKTLEKSTALAAIKPILE---NPNIHKIGQNIKFD-ESIFARHGIELQGVEFDTML 448
Query: 183 AAMLLNENESHSLKSLHSKYVRNEENAEVAKFNDLFKGIPFSLIPPDVAYMYAAYDPLQT 242
           + LN H++ L +Y+ +E A + + F+ IP + A YAA D
Sbjct: 449 LSYTLNSTGRHNMDDLAKRYLGHETIAFESLAGKGKSQLTFNQIPLEQATEYAAEDADVT 508
Query: 243 FELYEFQEQYLTPGTEQCEEYNLEKVSWVLHNIEMPLIKVLFDMEVYGVDLDQDKLAEIR 302
                                        +E+PL+ VL ME GV +D D L
                         EY
                   L
            +L +
Sbjct: 509 MKLQQALWLKLQEEPTLVELYK------TMELPLLHVLSRMERTGVLIDSDALFMQS 559
Query: 303 EQFTANMNEAEQEFQQLVSEWQPEIEELRQTNFQSYQKLEMDARGRVTVSISSPTQLAIL 362
                                                         +++S OL +
            + + + E++ L +
Query: 363 FYDIMGLKSPERDKPRG---TGESIVEH--FDNDISXXXXXXXXXXXXXXXXTTT-LDQHL 416
            +D + L ++ P+G T E ++E + +++
Sbjct: 593 LFDKLELPVLQKT-PKGAPSTNEEVLEELSYSHELPKILVKHRGLSKLKSTYTDKLPQMV 651
Query: 417 AKPDNRIHTTFKQYGAKTGRMSSENPNLQNIPSRGE-GAVVRQIFAASEGHYIIGSDYSQ 475
               R+HT++ Q TGR+SS +PNLQNIP R E G +RQ F A EG+ I+ +DYSQ
Sbjct: 652 NSQTGRVHTSYHQAVTATGRLSSSDPNLQNIPIRNEEGRHIRQAFIAREGYSIVAADYSQ 711
Query: 476 QEPRSLAELSGDESMRHAYEQNLDLYSVIGSKLYGVPYEBCLEFYPDGTTNKEGKLRRNS 535
E R +A LSGD+ + +A+ Q D++ ++++GV +B T+++ R +
Sbjct: 712 IELRIMAHLSGDQGLINAFSQGKDIHRSTAAEIFGVSLDE------VTSEQ----RRN 759
Query: 536 VKSVLLGLMYGRGANSIAEQMNVSVKEANKVIEDFFTEFPKVADYIIFVQQQAQDLGYVQ 595
           K++ GL+YG A ++ Q+ +S +A K ++ +F +P V ++ ++++A+ GYV+
```

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Sbjct: 760 AKAINFGLIYGMSAFGLSRQLGISRADAQKYMDLYFQRYPSVQQFMTDIREKAKAQGYVE 819
Query: 596 TATGRRRRLPDMS 608
           T GRR LPD++
Sbjct: 820 TLFGRRLYLPDIN 832
 Score = 46.9 bits (109), Expect = 5e-04
 Identities = 34/123 (27%), Positives = 66/123 (53%), Gaps = 16/123 (13%)
Query: 663 EIKDQAKAEGI------LIKDNGGKIADAQRQCLNSVIQGTAADMTKYAMIKV 709
                                   + N + A+R +N+ +QGTAAD+ K AMIK+
           +I+++AKA+G
Sbjct: 807 DIREKAKAQGYVETLFGRRLYLPDINSSNAMRRKGAERVAINAPMQGTAADIIKRAMIKL 866
Ouery: 710 HNDAELKELGFHLMIPVHDELLGEVPIKNAKRGAERLTEVMIEAAKDIISLPMKCDPSIV 769
                      +++ VHDEL+ EV +
                                            E++ + M EAA +++ +P+ + +
Sbjct: 867 -DEVIRHDPDIEMIMQVHDELVFEVRSEKVAFFREQIKQHM-EAAAELV-VPLIVEVGVG 923
Query: 770 ERW 772
           + W
Sbjct: 924 QNW 926
Query= sid|114825|lan|dp1ORF004 Phage dp1 ORF|40401-42440|3
         (679 letters)
>emb|CAB07981| (Z93946) hypothetical protein [bacteriophage Dp-1]
           Length = 532
 Score = 1011 bits (2585), Expect = 0.0
 Identities = 497/499 (99%), Positives = 498/499 (99%)
         MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLNG 60
Query: 1
           MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLNG
          MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLNG 60
Sbjct: 1
Query: 61 SSVHSSHPDYDTSGEEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL 120
           SSVHSSHPDYDTSGEEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL
Sbjct: 61 SSVHSSHPDYDTSGEEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL 120
Query: 121 DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGKNHTTSVSFT 180
           DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGKNHTTSVSFT
Sbjct: 121 DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGKNHTTSVSFT 180
Query: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT 240
           PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT
Sbjct: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT 240
Ouery: 241 SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF 300
           SAVROILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF
Sbjct: 241 SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF 300
Query: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTRQNPAIIQALRNAKVAPI 360
           NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTRQNPAIIQALRNAKVAPI
Sbjct: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTRQNPALIQALRNAKVAPI 360
Query: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISLMTNSSANLAGNYGPDKSYIV 420
           TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISL+TNSSANLAGNYGPDKSYIV
Sbjct: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISLLTNSSANLAGNYGPDKSYIV 420
Query: 421 KAKIQDRFTSTEFSATVATESVVLNYDKDGRLGVGKVVEQGKAGSIDAAGDIYAGGRQVQ 480
          KAKIQDRFTSTEFSATV TESVVLNYDKDGRLGVGKVVEQGKAGSIDAAGDIYAGGRQVQ
Sbjct: 421 KAKIQDRFTSTEFSATVPTESVVLNYDKDGRLGVGKVVEQGKAGSIDAAGDIYAGGRQVQ 480
Query: 481 QFQLTDNNGALNRGQYNDV 499
           OFOLTDNNGALNRGOYNDV
Sbjct: 481 QFQLTDNNGALNRGQYNDV 499
Query= sid|114827|lan|dp10RF006 Phage dp1 ORF|45296-46987|2
                                                                                (563 letters)
>gb|AAD18987| (AE001666) SWI/SNF family helicase_2 [Chlamydia pneumoniae]
 Score = 171 bits (429), Expect = 1e-41
 Identities = 150/522 (28%), Positives = 254/522 (47%), Gaps = 55/522 (10%)
```

```
SSNNFE-LPYKYFNNVIDALDEWELHIFGELDKDVQDYIDSRNRIASSSNEQFSFKTTPF 104
Query: 46
           S + FE LP + ++ + L E + I GE++ D QD
Sbjct: 659 SLDQFEALPVNF--SMSERLIEIQKQIRGEIEFDFQD------VPQQIQATLRSYQTEG 709
Query: 105 AHQVECFEYAQEHPCFLLGDEQGLGKTKQAIDIAVSRKASFKH--CLIVCCISGLKWNWA 162
            H +E + H +L D+ GLGKT QAI IAV++ K C ++ C + L +NW
           VHWLE--RLRKMHLNGILADDMGLGKTLQAI-IAVTQSKLEKGSGCSLIVCPTSLVYNWK 766
Sbjct: 710
Query: 163 KEVGIHSNESAHILGSRVTKDGKLVIDGV-SKRAEDLLGGHDEFFLITNIETLRDAVFIK 221
                                 LVIDGV S+R + L D IT+ L+ V
           +E
                + E
           EEFRKFNPEFR-----TLVIDGVPSQRRKQLTALADRDVAITSYNLLQKDV--- 812
Sbjct: 767
           YLNELTKSGEIGMVIIDEIHKCKNPSSKQGASIQKLQSYYKMGLTGTPLMNNPIDVFNVM 281
Ouerv: 222
                     V++DE H KN +++ S++ +QS +++ LTGTP+ N+ +++++
           ---ELYKSFRFDYVVLDEAHHIKNRTTRNAKSVKMIQSDHRLILTGTPIENSLEELWSLF 869
Shict: 813
           KWLGAEHHTLTQFKERYCIVDQFNQITGYR----NLAELRELVNDYMLRRTKEEVL-DL 335
                L +R+ V ++ + Y N+ L++ V+ ++LRR KE+VL DL
           DFLMPG---LLSSYDRF--VGKYIRTGNYMGNKADNMVALKKKVSPFILRRMKEDVLKDL 924
           PEKIRVTEYVDMNSKQSKIY------KEVLTKLVQEIDKVKLMPNPLAETIRLRQATGN 388
P + + + Q ++Y K+ L++LV++ ++ + LA RL+Q +
Query: 336
           PPVSEILYHCHLTESQKELYQSYAASAKQELSRLVKQEGFERIHIHVLATLTRLKQICCH 984
Sbict: 925
           PSILTTQDVK---SCKFERCIEIVEECIQQGKSCVIFSNWEKVIEPLAKIL-SKTVKCNL 444
                  + S K++ ++++ + G V+FS + K++ + K L S+ +
Sbjct: 985 PAIFAKDAPEPGDSAKYDMLMDLLSSLVDSGHKTVVFSQYTKMLGIIKKDLESRGIPFVY 1044
Query: 445 VTGETADKFNEIEEFMNHRKASVILGTIGALGTGFTLTKADTVIFLDSPWTRAEKDQAED 504
            Sbjct: 1045 LDGSTKNRLDLVNQFNEDPSLLVFLISLKAGGTGLNLVGADTVIHYDMWWNPAVENQATD 1104
Query: 505 RCHRIGAKSSVTIYTLVAKGTVDERIEDLIERKGELADYIVD 546
           R HRIG SV+ Y LV T++E+I L RK L
Sbjct: 1105 RVHRIGQSRSVSSYKLVTLNTIEEKILTLQNRKKSLVKKVIN 1146
Query= sid|114828|lan|dp10RF007 Phage dp1 ORF|22230-23621|3
         (463 letters)
>gi|2444105 (U88974) ORF26 (Streptococcus thermophilus temperate bacteriophage
          01205}
          Length = 411
Score = 88.9 bits (217), Expect = 7e-17
Identities = 80/315 (25%), Positives = 133/315 (41%), Gaps = 48/315 (15%)
Query: 139 QGVTLAGIFCDEVALMPESFVNQATGRCSVTGSKMWFSCNPANPNHYFKKNWIDKQVEKR 198
          +G T G + +E +L E + RCS G+++ + NP NPNH+ +++I K + -
Sbjct: 121 RGFTAFGAYVNEASLANELVFKEIISRCSGDGARVVWDSNPDNPNHWLNRDYIGKN-DGK 179
Query: 199 ILYLHFTMDDNPSLT----DSIKRRYEKMYAGVFRKRFILGLWVTADGLVYSMFNEEQHV 254
I+ F+DDN L+ DSIK K G F R ILGLW A+G +Y+ ++ HV
Sbjct: 180 IIDFSFKLDDNTFLSKRYIDSIKAATPK---GKFYDRDILGLWTVAEGAIYADYDSKIHV 236
Query: 255 KKLNIEFDRLFVAGDFGIYNATTFGLYGFSKRHKRYHLIESYYHSGREAEEQLTEADVNS 314
               E R F D+G + + + G
                                         ++L++
Sbjct: 237 VDELPEMKRYFGGIDWGYTHYGSIVIVG-EGVDNNFYLVDGVAAQFKEIDWWVEQA---- 291
Query: 315 NIQFSSVLQKTTKEYANDLVDMIRGKQIEYIILDPSASAMIVELQKHPYIAR---KNIPI 371
                                                   + + ++AR
                  +K T Y N
Sbjct: 292 -----RKLTGIYGN------IPFYADSARPEHVARFENEGFDI 323
Query: 372 IPARNDVTLGISFHAELLAENRFTLDPSNT-HDIDEYYAYSWDSKASQTGEDRVIKEHDH 430
          + A V GI A+L E + +
                                          DE Y Y W ++ +D +KE D
Sbjct: 324 MNANKSVIAGIELIAKLFKEKKLYVKRGFVPRFFDEIYQYRWKENST---KDEPLKEFDD 380
Query: 431 CMDRNRYACLTDALI 445
                                                                              ____
           +D RYA +D +I
Sbjct: 381 VLDSVRYAIYSDYVI 395
Query= sid|114829|lan|dplORF008 Phage dpl ORF|49624-50961|1
        (445 letters)
```

>gb|AAD19901| (AF100420) DnaB replication fork helicase [Thermus aquaticus]

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Length = 444
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Score = 67.5 bits (162), Expect = 2e-10
 Identities = 69/248 (27%), Positives = 111/248 (43%), Gaps = 14/248 (5%)
Query: 147 GERLGISTGFEXXXXXXXXXXXXXXXXIVIMARPGQGKS-WTIDKMLATAWKNGHDVLLYS 205
                                    I I ARP GK+ + +
                                                         AKG V+YS
           GE G+ TGF+
Sbjct: 178 GEVAGVRTGFKELDQLIGTLGPGSLNI-IAARPAMGKTAFALTIAQNAALKEGVGVGIYS 236
Query: 206 GEMSEMQVGARIDTILSNVSINSITKGIWNDHQFEKYEDHIQAMTEAENSLVVVTPFMIG 265
            EM Q+ R+ +++N+ G D F + D
                                                    ++EA
Sbjct: 237 LEMPAAQLTLRMMCSEARIDMNRVRLGQLTDRDFSRLVDVASRLSEAP-IYIDDTPDLTL 295
Query: 266 GKNLTPAILDSMISKYRPSVVGIDQLSLMS--ESYPSREQKRIQYANITMDLYKISAKYG 323
+ A ++S+ + ++ ID L LMS S S E ++ + A I+ L ++ + G
Sbjct: 296 ME--VRARARRLVSQNQVGLIIIDYLQLMSGPGSGKSGENRQQEIAAISRGLKALARELG 353
Query: 324 IPIVLNVQAGRSAKTEGAESMELEHIAESDGVGQNASRVIAMKRD-----EKSGILEL 376
IPI+ Q R+ + + L + ES + Q+A V+ + RD EK+GI E+
Sbjct: 354 IPIIALSQLSRAVEARPNKRPMLSDLRESGSIEQDADLVMFIYRDEYYNPHSEKAGIAEI 413
Query: 377 SVVKNRYG 384
            VKRG
Sbjct: 414 IVGKQRNG 421
Query= sid|114831|lan|dp10RF010 Phage dp1 ORF|8699-9859|2
         (386 letters)
>gi|2760912 (AF037258) RecA protein [Chlorobium tepidum]
          Length = 346
 Score = 133 bits (331), Expect = 2e-30
 Identities = 99/340 (29%), Positives = 164/340 (48%), Gaps = 66/340 (19%)
GGLPR RV E +GPESSGKTT AL + AQ
Sbjct: 67 GGLPRGRVTEIYGPESSGKTTLALHAIAEAQ-----KNG 100
Query: 104 AVKELEMQLDSLQEPLKIVYLDLENTLDTEWAKKIGVDVDNIWIVRPEMNSAEEILQYVL 163
                            +D E+ D +A+K+GVD++ + + +PE S E+ L V
Sbjct: 101 GIAAL-----VDAEHAFDPTYARKLGVDINALLVSQPE--SGEQALSIVE 143
Query: 164 DIFETGEVGLVVLDSLPYMVSQNLIDEELTKKAYAGISAPLTEFSRKVTPLLTRYNAIFL 223
                                             + +++ RK+T +++ L
           + +G V ++V+DS+ +V Q ++ E+
Sbjct: 144 TLVRSGAVDIIVIDSVAALVPQAELEGEMGDSVVGLQARLMSQALRKLTGAISKSSSVCL 203
Query: 224 GINQIREDMNSQYNA-YSTPGGKMWKHACAVRLKFRKGDYLDENGASLTRTARNPAGNVV 282
           INQ+R+ + Y + +T GGK K +VRL RK + ++G L
Sbjct: 204 FINQLRDKIGVMYGSPETTTGGKALKFYSSVRLDIRKIAQI-KDGEELV------GNRT 255
Query: 283 ESFVEKTKAFKPDRKLVSYTLSYHDGIQIENDLVDVAVEFGVIQKAGAWFSIVDLETGEI 342
           + V K K P K + + Y +GI + +L+D+AVEFG+I+K+GAWFS
Sbjct: 256 KVKVVKNKV-APPFKTAEFDILYGEGISVLGELIDLAVEFGIIKKSGAWFSYGTEKLG-- 312
Query: 343 MTDEDEEPLKFQGKANLVRRFKEDDYLFDMVMTAVHEIIT 382
                     QG+ N+ + KED+ L + + V +++T
Sbjct: 313 -----QGRENVKKLLKEDETLRNTIRQQVRDMLT 341
Query= sid | 114832 | lan | dp1ORF011 Phage dp1 ORF | 28017-29096 | 3
         (359 letters)
>gi|2444110 (U88974) ORF31 (Streptococcus thermophilus temperate bacteriophage
          01205]
          Length = 348
 Score = 187 bits (469), Expect = 1e-46
 Identities = 118/358 (32%), Positives = 187/358 (51%), Gaps = 21/358 (5%)
Query: 3 IYDYINAGEIASYIQALPSNALQYLGPTLFPNAQQTGTDISWLKGANNLPVTIQPSNYDA 62 IYD + A IA Y AL N LG ++FP +Q GT +S++KGA+ V ++ +D
Sbjct: 4 IYDKVTASNIAGYFNALQENVSSTLGESIFPARKQLGTKLSYIKGASGQSVALKAAAFDT 63
Query: 63 KASLRERAGFSKQATEMAFFRESMRLGEKDRQNLQMLLNQSSA-LAQPLITQLYNDTKNL 121
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+M FF+E+M + E DRO L ++ + +A L
Sbjct: 64 NVTIRDRVSAEMHDEQMPFFKEAMLVKENDRQQLNLVKDSGNAVLVNTIVAGIFNDNLTL 123
Query: 122 VDGVEAQAEYMRMQLLQYGKFTVKSTNSEAQYTYDYNMDAKQQYAVTKKWTNPAESDPIA 181
V+G A+ B MRMQ+L GK S Y D K+Q V+K W P + P+A
Sbjct: 124 VNGARARLEAMRMQVLATGKIAFTSDGVNKDIDYGVKPDHKKQ--VSKSWAEPG-ATPLA 180
Query: 182 DILAAMDDIENRTGVRPTRMVLNRNTYNQMTKSDSIKKAL-AIGVQGSWENFLLLASDAE 240
D+ A+ + G+ P R V+N T+ + K+ S K + + GS + + + E
Sbjct: 181 DLEDAI-ETARELGLNPERAVMNAKTFGLIRKAASTVKVIKPLAGDGS----AVTKAELE 235
Query: 241 KFIAEKTGLQIAVYSKKIAQFADADKLPDVGNIRQFNLIDDGKVVLLPPDAVGHTWYGTT 300
            +IA+ G+ I + +
                                      DG + +F DG + L+P +G+T +GTT
Sbict: 236 NYIADNFGVSIVLENGTYRN------DKGEVSKF--YPDGHLTLIPNGPLGNTVFGTT 285
Ouery: 301 PEAFDLASGGT-DAQVOVLSGGPTVTTYLEKHPVNIATVVSAVMIPSFEGIDYVGVLT 357
           PE DL + T +A+V+++ G VTT PVN+ T VS V +PSFE +D V +LT
Sbict: 286 PEESDLFADNTVNAEVEIVDNGIAVTTTKTTDPVNVOTKVSMVALPSFERLDDVYMLT 343
Query= sid | 114834 | lan | dp10RF013 Phage dp1 ORF | 10215-11240 | 3
         (341 letters)
>sp|P09122|DP3X BACSU DNA POLYMERASE III SUBUNITS GAMMA AND TAU
           Length = 563
 Score = 182 bits (458), Expect = 2e-45
 Identities = 118/353 (33%), Positives = 176/353 (49%), Gaps = 31/353 (8%)
           YRPOTFEEVVAQEYVKEILLNQLQNGAIKHGYLFCXXXXXXXXXXXIFAKDVN----- 60
Ouerv: 7
           +RPO FE+VV OE++ + L N L H YLF
                                                         +IFAK VN
Sbjct: 10 FRPORFEDVVGQEHITKTLONALLOKKFSHAYLFSGPRGTGKTSAAKIFAKAVNCEHAPV 69
Query: 61 ------KGL----GSPIEIDAASNNGVENVRNIIEDSRYKSMDSEFKVYIIDEVH 105
                               IEIDAASNNGV+ +R+I + ++
Sbjct: 70 DEPCNECAACKGITNGSISDVIEIDAASNNGVDEIRDIRDKVKFAPSAVTYKVYIIDEVH 129
Query: 106 MLSTGAFNALLKTLEEPSSGTVFILCTTDPQKIPDTILSRVQRFDFTRIDNDDIVNOLOF 165
           MLS GAFNALLKTLEEP +FIL TT+P KIP TI+SR QRFDF RI + IV ++
Sbjct: 130 MLSIGAFNALLKTLEEPPEHCIFILATTEPHKIPLTIISRCQRFDFKRITSQAIVGRMNK 189
Query: 166 IIESENEEGAGYSYERDALSFIGKLANGGMRDSITRLEKVLDYSHHVDMEAVSNAL---G 222
                       E +L I A+GGMRD+++ L++ + +S D+ V +AL
Sbjct: 190 IVDAEQ-----LQVEEGSLEIIASAAHGGMRDALSLLDQAISFSG--DILKVEDALLITG 242
Query: 223 VPDYETFASLVEAIANYDGSKCLEIVNDFHYSGKDLKLVTRNFTDFLLEVCKYWLVRDIS 282
                    L +++ + + S LE +N+ GKD + + + ++ Y
Sbict: 243 AVSQLYIGKLAKSLHDKNVSDALETLNELLQQGKDPAKLIEDMIFYFRDMLLYKTAPGLE 302
Query: 283 ITQLPAHFESKLEQFCEAFQYPTLLWMLEEMNELAGVVKWEPNAKPIIETKLL 335
                                 L M++ +N+ +KW + + E ++
                       + E
Sbjct: 303 GVLEKVKVDETFRELSEQIPAQALYEMIDILNKSHQEMKWTNHPRIFFEVAVV 355
Query= sid|114835|lan|dp10RF014 Phage dp1 ORF|50961-51974|3
         (337 letters)
>sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64227 DNA primase (dnaE) homolog
           MG250 - Mycoplasma genitalium (SGC3) >gi|3844848
           (U39704) DNA primase (dnaE) [Mycoplasma genitalium]
           Length = 607
 Score = 57.0 bits (135), Expect = 2e-07
 Identities = 53/190 (27%), Positives = 89/190 (45%), Gaps = 17/190 (8%)
Query: 146 EELDKYRFIHP------YMYERKLTDELIEMFDVGYDK--LHDCITFPVRNLKGETVFF 196
E +++Y FI+P Y++ K + + FD K + I P+ + G V F
Sbjct: 170 ESMERYPFINPKIKPSELYLFS-KTNQQGLGFFDFNTKKATFQNQIMIPIHDFNGNPVGF 228
                                                                                 Query: 197 NRRSVRSKFHQYGEDDPKTEFLYGQYELVAFRDYFEKPISQVFVTESVINCLTLWSMKIP 256 -
                             EF + + EL+
                                              K ++Q+F+ E
Sbjct: 229 SARSVDNINKLKYKNSADHEF-FKKGELLFNFHRLNKNLNQLFIVEGYFDVFTLTNSKFE 287
Query: 257 AVALMGVGGGN-QINLLKR--LPYRNIVLALDPDNAGQTAQEKLYRQLKRSK-VVRFLNY 312
          AVALMG+ + QI +K + +VLALD D +GQ A L +L + +V + +
Sbjct: 288 AVALMGLALNDVQIKAIKAHFKELQTLVLALDNDASGQNAVFSLIEKLNNNNFIVEIVQW 347
```

Query: 313 PKEFYDNKWD 322

+ D WD

<u>\_\_\_\_</u>

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Sbjct: 348 EHNYKD--WD 355
Query= sid | 114837 | lan | dp10RF016 Phage dp1 ORF | 43413-44303 | 3
         (296 letters)
>emb|CAB07986| (293946) N-acetylmuramoyl-L-alanine amidase [bacteriophage Dp-1]
           Length = 296
 Score = 661 bits (1686), Expect = 0.0
 Identities = 296/296 (100%), Positives = 296/296 (100%)
          MGVDIEKGVAWMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60
           MGVDIEKGVAWMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH
Sbjct: 1
          MGVDIEKGVAWMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60
Query: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSDNIIHCNYAYDGIS 120
           AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSDNIIHCNYAYDGIS
Sbjct: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSDNIIHCNYAYDGIS 120
Query: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180
           VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE
Sbjct: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180
Query: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYYFNRDGSMVTGW 240
           ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYYFNRDGSMVTGW
Sbjct: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYYFNRDGSMVTGW 240
Query: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV 296
          IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV
Sbjct: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV 296
Query= sid|114841|lan|dp1ORF020 Phage dp1 ORF|1864-2658|1
         (264 letters)
>emb|CAB13247| (Z99111) similar to coenzyme PQQ synthesis [Bacillus subtilis]
          Length = 243
Score = 217 bits (548), Expect = 5e-56
Identities = 117/248 (47%), Positives = 163/248 (65%), Gaps = 15/248 (6%)
Ouery: 23 MPIMEIFGPTIOGEGMVIGOKTIFIRTGGCDYHCNWCDSAFTWNGTTEPE--YITGKEAA 80
           +P++EIFGPTIQGEGMVIGQKT+F+RT GCDY C+WCDSAFTW+G+ + + ++T +E
```

Query: 141 KEVSDITISPKPPSSGMRTNMKILEAIVDRM--NDENLDWSFKIVIFDENDLAYARDMFK 198
+ D+TISPKPPSS M TN + L+ I+ + ND S K+VIF++ DL +A+ + K
Sbjct: 119 TLIDDLTISPKPPSSKMVTNFQKLDHILTSLQENDRQHAVSLKVVIFNDEDLEFAKTVHK 178

Query: 199 TFEGKLRPVNYLSVGNANAY--EEGKISDRLLEKLGWLWDKVYEDPAFNNVRPLPQLHTL 256
+ G YL VGN + + ++ + LL K L DKV D N VR LPQLHTL
Sbjct: 179 RYPG---IPFYLQVGNDDVHTTDDQSLIAHLLGKYEALVDKVAVDAELNLVRVLPQLHTL 235

Query: 257 VYDNKRGV 264

Query: 81 SRILKLAFNDKGEQICNHVTLTGGNPALINEPMAKMISILKEHGFKFGLETQGTRFQEWF 140

Sbjct: 65 AEL----KDIGGDAFSHVTISGGNPALLKQ-LDAFIELLKENNIRAALETQGTVYQDWF 118

IPVLEIFGPTIQGEGMVIGQKTMFVRTAGCDYSCSWCDSAFTWDGSAKKDIRWMTAEEIF 64

+HVT++GGNPAL+ + + I +LKE+ + LETQGT +Q+WF

++ NKRGV Sbjct: 236 LWGNKRGV 243

DG

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Query= sid|114842|lan|dplORF021 Phage dp1 ORF|2504-3295|2
>8P|P19465|GCH1_BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >gi|98411|pir||A38256 GTP
          cyclohydrolase I (EC 3.5.4.16) - Bacillus subtilis
           >gi|143231 (M37320) regulatory protein (Bacillus
           subtilis] >gi|143799 (M80245) MtrA [Bacillus subtilis]
           >gi|2634696|emb|CAB14194| (Z99115) GTP cyclohydrolase I
           [Bacillus subtilis]
          Length = 190
Score = 208 bits (523), Expect = 4e-53
Identities = 103/185 (55%), Positives = 133/185 (71%), Gaps = 1/185 (0%)
Query: 80 VTLDNTEAAVQRLFGLLGEDAERDGLQDTPFRFVKALAEHTVGYREDPKLHLEKTFDVDH 139
          V + E AV+++ +GED R+GL DTP R K AE G EDPK H + F +H
          VNKEQIEQAVRQILEAIGEDPNREGLLDTPKRVAKMYAEVFSGLNEDPKEHFQTIFGENH 63
Query: 140 EDLVLVKDIPFNSLCEHHLAPFVGKVHIAYIPKD-KITGLSKFGRVVEGYAKRLQVQERL 198
           E+LVLVKDI F+S+CEHHL PF GK H+AYIP+ K+TGLSK R VE AKR Q+QER+
Sbjct: 64 EELVLVKDIAFHSMCEHHLVPFYGKAHVAYIPRGGKVTGLSKLARAVEAVAKRPQLQERI 123
Query: 199 TQQIADAIQEVLNPQAVAVIVEAEHTCMSGRGIKKHGATTVTSTMRGLFQDDASARAELL 258
          T IA++I E L+P V V+VEAEH CM+ RG++K GA TVTS +RG+F+DDA+ARAE+L
Sbjct: 124 TSTIAESIVETLDPHGVMVVVEAEHMCMTMRGVRKPGAKTVTSAVRGVFKDDAAARAEVL 183
Query: 259 QLIKK 263
           + IK+
Sbjct: 184 EHIKR 188
Query= sid | 114843 | lan | dp10RF022 Phage dp1 ORF | 30896-31675 | 2
         (259 letters)
>gi[2347102 (U77367) internalin [Listeria monocytogenes]
          Length = 821
 Score = 55.0 bits (130), Expect = 5e-07
 Identities = 44/149 (29%), Positives = 63/149 (41%), Gaps = 13/149 (8%)
Query: 119 FRMNIYVPNYVG--DSIVNYVKITLNNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPV 176
F + VPN + D + + NN T AP L Y PE +K + K +
Sbjct: 383 FSKTLSVPNNITSIDGTLIAPETISNNGTYDAPNLKWSLPNYLPE--VKYTFSQKIPIGT 440
Query: 177 KSMDYVAQLPAVLR-----RVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW 231
            + +Y + L+ +VTF++ G T + V E + P+P PT G F GW
Sbjct: 441 GTSNYSGFITQPLKELLDYKVTFNVEGNTSEVETVTEE---NLIPEPTSPTKQGYTFDGW 497
Query: 232 -KVEGESTIWDFDNHMMPDRDVKLVAQFA 259
             E T WDF MP D+ L A F+
Sbjct: 498 YDAETGGTKWDFTTGQMPANDLTLYAHFS 526
 Score = 43.4 bits (100), Expect = 0.002
 Identities = 47/195 (24%), Positives = 73/195 (37%), Gaps = 12/195 (6%)
Query: 72 YDLTFKDNTFDPEIMALIEGGTVRQQGGTIAGYDT-PMLAQGASNMKPFRMNIYVPNY-- 128
                + T + +G + GG + T M A + P +N Y N+
Sbjct: 547 YDALLNEPTTPTKQGYTFDGWYDAETGGNKWDFKTMKMPANDVAFYAHFTINNYQANFDI 606
Query: 129 ---VGDSIVNYVKITLMNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPVKSMDYVAQL 185
              V + + Y + T G + + A
                                                K TK +P
Sbjct: 607 DGEVKNETIAYDTLLNEPTTPTKQGYTFDGWYDAETGGTKWDFKTKE-MPANDVTLYAHF 665
Query: 186 PAVLRRVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW-KVEGESTIWDFDN 244
+ FD++G T + V +A + P+P P+ TG +GW E T WDF
Sbjct: 666 TINNYQANFDIDGAV-TEEVVNYDA---LIPEPTSPSKTGFTLEGWYDAEVGGTKWDFKT 721
Query: 245 HMMPDRDVKLVAQFA 259
                                                                              MP D+ L A F+
Sbict: 722 MKMPANDITLYAHFS 736
 Score = 38.3 bits (87), Expect = 0.057
 Identities = 42/169 (24%), Positives = 59/169 (34%), Gaps = 10/169 (5%)
```

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Query: 96 QQGGTIAGYDT-PMLAQGASNMKPFRMNIYVPNYVGDSIVNYVKIT----LNNCTGKAPG 150
                                                           LN T
            + GGT + T M A + F +N Y N+ D +V +
Sbjct: 501 ETGGTKWDFTTGQMPANDLTLYAHFSVNSYQANFDIDGVVTNEAVVYDALLNEPTTPTKQ 560
Query: 151 LSIGKEFYAPEFNIKAREATKAGLPVKSMDYVAQLPAVLRRVTFDLNGGTGTADAVRVEA 210
                +Y E + +P + + A + FD++G
Sbjct: 561 GYTFDGWYDAETGGNKWDFKTMKMPANDVAFYAHFTINNYQANFDIDGEVKNETI----A 616
Query: 211 GKKISPKPVDPTLTGKAFKGW-KVEGESTIWDFDNHMMPDRDVKLVAQF 258
+ +P PT G F GW E T WDF MP DV L A F
Sbjct: 617 YDTLLNEPTTPTKQGYTFDGWYDAETGGTKWDFKTKEMPANDVTLYAHF 665
Ouery= sid|114850|lan|dp1ORF029 Phage dp1 ORF|662-1348|2
         (228 letters)
>gi|2650185 (AE001074) succinoglycan biosynthesis regulator (exsB)
           [Archaeoglobus fulgidus]
           Length = 239
 Score = 119 bits (295), Expect = 2e-26
 Identities = 79/224 (35%), Positives = 113/224 (50%), Gaps = 11/224 (4%)
         MKSVVLLSGGVDSATCLAIEVDKWGSKNVHAIAFNYGQKHEAELENAANVAMFYGVKFTI 60
           MK+V+LLSGG+DS+T L +D G VHA+ F YGQKH E+E+A VA
          MKAVMLLSGGIDSSTLLYYLLD--GGYEVHALTFFYGQKHSKEIESAEKVAKAAKVRHLK 58
Sbjct: 1
Query: 61 LEIDSKIYXXXXXLLQGKGEISHGKSYAEILAEKEVVDTYVPFRNGLMLSQXXXXXXXX 120
           ++I S I+ L G+ E+ Y+E + + T VP RN ++LS
Sbict: 59 VDI-STIHDLISYGALTGEEEVPKA-FYSEEVQRR----TIVPNRNMILLS--IAAGYAV 110
Query: 121 XXXXXXXXXXXXXXXXXXXXXXXPDCTPEFYNSMSNAMEYGT-GGKVTLVAPLLTLTKAQVVKW 179
                              PDC EF ++ A+
                                                  V + AP + +TKA +V+
Sbjct: 111 KIGAKEVHYAAHLSDYSIYPDCRKEFVKALDTAVYLANIWTPVEVRAPFVDMTKADIVRL 170
Query: 180 GIDLDVPYFLTRSCYESDAESCGTCATCIDRKKAFEENGMTDPI 223
           G+ L VPY LT SCYE C +C TC++R +AF NG+ DP+
Sbjct: 171 GLKLGVPYELTWSCYEGGDRPCLSCGTCLERTEAFLANGVKDPL 214
Query= sid | 114855 | lan | dplORF034 Phage dpl ORF | 131-652 | 2
         (173 letters)
>emb|CAB13248| (299111) similar to hypothetical proteins [Bacillus subtilis]
          Length = 165
 Score = 220 bits (556), Expect = 4e-57
 Identities = 103/139 (74%), Positives = 117/139 (84%)
Query: 5 TTRTDAELTGVTLLGNQDTKYDYDYNPDVLETFPNKHPENNYLVTFDGYEFTSLCPKTGQ 64
           TTR ++EL GVTLLGNQ T Y ++Y PDVLE+FPNKH +Y V F+ EFTSLCPKTGQ
Sbjct: 2 TTRKESELEGVTLLGNQGTNYLFEYAPDVLESFPNKHVNRDYFVKFNCPEFTSLCPKTGQ 61
Query: 65 PDFANVFISYIPNEKMVESKSLKLYLFSFRNHGDFHEDCMNIILNDLYELMEPKYIEVMG 124
           PDFA ++ISYIP+EKMVESKSLKLYLFSFRNHGDFHEDCMNII+NDL ELM+P+YIEV G
Sbjct: 62 PDFATIYISYIPDEKMVESKSLKLYLFSFRNHGDFHEDCMNIIMNDLIELMDPRYIEVWG 121
Query: 125 LFTPRGGISIYPFVNKVNP 143
           FTPRGGISI P+ N P
Sbjct: 122 KFTPRGGISIDPYTNYGKP 140
Query= sid|114857|lan|dp10RF036 Phage dp1 ORF|48808-49362|1
         (184 letters)
>gi|1353529 (U38906) ORF12 [Bacteriophage rlt]
          Length = 296
                                                                             - -
 Score = 53.5 bits (126), Expect = 1e-06
 Identities = 42/149 (28%), Positives = 70/149 (46%), Gaps = 9/149 (6%)
Query: 34 IASNTVGNGKTSWAVRLLQRYLAETALDGRIVEKGMFVVSAQLLTEFGDYNYFQTMQEFL 93
+ S G GK+ A+ +L+ L T L ++ V + F + + F + F + F + Sbjct: 155 VVSGPAGTGKSHLAMSILKDCLQHTDLT--VIFASWSEVLHLIKDSFDNKDSFYSTEYFM 212
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Query: 94 ERFERLKTCELLVIDEIGGGSLTKASYPYLYDLVNYRVDNNLSTIYTTNYTDDEIIDLLG 153
           E F + +LLVID+IG +T+ S L ++++ R
                                                     TI TTN DEI
Sbjct: 213 EVF---RNTDLLVIDDIGSEKITEWSMSLLTEVLDART----KTIITTNLKSDEIRKKYH 265
Query: 154 QRLYSRIYDTSVVLDFQASNVRGLEVSEI 182
            R YSR++ P N++ VS++
Sbjct: 266 NRTYSRLFRGIGKKAFNFENIKDKRVSQL 294
Query= sid|114859|lan|dp1ORF038 Phage dp1 ORF|1350-1871|3
         (173 letters)
>sp|P44123|YB90 HAEIN HYPOTHETICAL PROTEIN HI1190 >gi|1074675|pir||F64021 hypothetical
           protein HI1190 - Haemophilus influenzae (strain Rd KW20)
           >gi|1574117 (U32798) 6-pyruvoyl tetrahydrobiopterin
           synthase, putative [Haemophilus influenzae Rd]
           Length = 141
 Score = 100 bits (247), Expect = 6e-21
 Identities = 59/143 (41%), Positives = 83/143 (57%), Gaps = 10/143 (6%)
Query: 2 RVSKTLTFDAAHQLVGHFGKCANLHGHTYKVEISLAGGTYDHGSSQGMVVDFYHVKKIA- 60
+SK +FD AH L GH GKC NLHGHTYK+++ ++G Y G+ + MV+DF +K I
Sbjct: 3 KISKEFSFDMAHLLDGHDGKCQNLHGHTYKLQVEISGDLYKSGAKKAMVIDFSDLKSIVK 62
Query: 61 GTFIDRLDHAVLL-QGNEP----IALANAVDTKRVLFGFRTTAENMSRFLTWTLTELMWK 115
+D +DHA + Q NE L +++K FRTTAE ++RF+ L +
Sbjct: 63 KVILDPMDHAFIYDQTNERESQIATLLQKLNSKTFGVPFRTTAEEIARFIFNRLKH--DE 120
Query: 116 HARIDSIKLWETPTGCAECTYYE 138
              I SI+LWETPT + C Y E
Sbjct: 121 QLSISSIRLWETPT--SFCEYQE 141
Query= sid|114860|lan|dp10RF039 Phage dp1 ORF|3306-3803|3
         (165 letters)
>emb|CAA68244| (X99978) ORF7; hydophobic protein [Lactobacillus plantarum]
          Length = 168
 Score = 64.4 bits (154), Expect = 5e-10
 Identities = 49/156 (31%), Positives = 84/156 (53%), Gaps = 9/156 (5%)
          WLVRTALIAALYVTLTVAFSAISY--GPIQFRVSEALILLPLWNHRWTPGIVLGTIIANF 65
          W++ AL+AA+YV L + +A S G IQFRVSE L L ++N ++ GIV G I+ +
          WIIN-ALVAAMYVVLCLGPAAFSLASGAIQFRVSEGLNHLAVFNRKYIWGIVAGVILFDA 67
Sbict: 9
Query: 66 FSP-LGLIDVLFGSLATFLGXXXXXXXXXXXXXXXICPVLA----NAYLIALELRIVY 120
           F P L++VLFG + L
Sbjct: 68 FGPGASLLNVLFGGGQSLLALLVLTWLAPKLKTVWQRMLLNIALFTVSMFMIALMITMMS 127
Query: 121 S-LPFWESVIYVGISEAIIVLISYFLISTLAKNNHF 155
          Sbjct: 128 SGVAFWPTYLTTALSELIIMSITAPIMYSLDRVLHF 163
Query= sid | 114862 | lan | dp1ORF041 Phage dp1 ORF | 8208-8699 | 3
         (163 letters)
>gi|2522313 (AF012906) dUTPase homolog [Bacillus subtilis]
          >gi|2634394|emb|CAB13893| (Z99114) similar to
          deoxyuridine 5'-triphosphate nucleotidohydrolase
           [Bacillus subtilis] >gi|3025643 (AF020713) putative
          dUTPase [Bacteriophage SPBc2]
          Length = 142
 Score = 108 bits (267), Expect = 2e-23
 Identities = 65/160 (40%), Positives = 83/160 (51%), Gaps = 25/160 (15%)
                                                                                ____
         VDVKMIDPKLDRLKYT--GDWVDVRISSITKIDADSADVSRCRKVLQKAQVYSVAAGECI 62
           + +K +D R+ GDW+D+R + I D
Sbjct: 3 IKIKYLDETQTRINKMEQGDWIDLRAAEDVAIKKDEFKL----------- 41
Query: 63 KIAHGFALELPKGYEAILHPRSSLFKKTGLIFVSS-GVIDEGYKGDTDEWFSVWYATRDA 121
           + G A+ELP+GYEA + PRSS +K G+I +S GVIDE YKGD D WF YA RD
Sbjct: 42 -VPLGVAMELPEGYEAHVVPRSSTYKNFGVIQTNSMGVIDESYKGDNDFWFFPAYALRDT 100
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WO 00/32825 PCT/IB99/02040

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Query: 122 DIFYDQRIAQFRIQEKQPAIKFNFVESLGNAARGGHGSTG 161
                RI QFRI +K PA+ V+ LGN RGGHGSTG
Sbjct: 101 KIKKGDRICOFRIMKKMPAVDLIEVDRLGNGDRGGHGSTG 140
Query= sid|114867|lan|dplORF046 Phage dp1 ORF|42774-43202|3
         (142 letters)
>emb|CAB07984| (293946) hypothetical protein [bacteriophage Dp-1]
          Length = 142
 Score = 287 bits (728), Expect = 2e-77
 Identities = 142/142 (100%), Positives = 142/142 (100%)
          MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNKAKSVLEDISTTLSTLKQQVDGIDQ 60
           MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNKAKSVLEDISTTLSTLKQQVDGIDQ
          MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNKAKSVLEDISTTLSTLKQQVDGIDQ 60
Sbjct: 1
Query: 61 TTVAINHQNDVIQDGTRKIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120
           TTVAINHQNDVIQDGTRKIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE
Sbjct: 61 TTVAINHQNDVIQDGTRKIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120
Query: 121 VEALYEKYKKLPIREEDLDETI 142
           VEALYEKYKKLPIREEDLDETI
Sbjct: 121 VEALYEKYKKLPIREEDLDETI 142
Query= sid|114901|lan|dplORF080 Phage dp1 ORF|42490-42759|1
         (89 letters)
 >emb|CAB07983| (Z93946) hypothetical protein [bacteriophage Dp-1]
          Length = 124
  Score = 147 bits (367), Expect = 1e-35
 Identities = 75/75 (100%), Positives = 75/75 (100%)
 Query: 1 MLNLTKSRQIVAEFTIGQGAEKKLVKTTIVNIDANAVSTVSETLHDPDLYAANRRELRAD 60
          MLNLTKSRQIVAEFTIGQGAEKKLVKTTIVNIDANAVSTVSETLHDPDLYAANRRELRAD
 Sbjct: 1 MLNLTKSRQIVAEFTIGQGAEKKLVKTTIVNIDANAVSTVSETLHDPDLYAANRRELRAD 60
 Query: 61 EQKLRETRYAIEDEI 75
           EQKLRETRYALEDE1
 Sbjct: 61 EQKLRETRYAIEDBI 75
 Query= sid|114912|lan|dp10RF091 Phage dp1 ORF|43189-43413|1
          (74 letters)
 >emb|CAB07985| (Z93946) holin [bacteriophage Dp-1]
           Length = 74
  Score = 63.2 bits (151), Expect = 2e-10
Identities = 34/74 (45%), Positives = 34/74 (45%)
 VLGVSSR
                                          YOFD
 Sbjct: 1 MKLSNEQYDVAKNVVTVVVPAAIALITGLGALYQFDTTAITGTIALLATFAGTVLGVSSR 60
           MKLSNEQYD
 Query: 61 NYQKEQEAQNNEVE 74
           NYQKEQEAQNNEVE
  Sbjct: 61 NYQKEQEAQNNEVE 74
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### Condensed listing of homology information from above

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Phage: dpl
Database: nr
Program: Blastp
Query= sid|114822|lan|dp10RF001 Phage dp1 ORF|36698-40390|2
         (1230 letters)
gi|2444124 (U88974) ORF45 (Streptococcus thermophilus temperate ...
                                                                        467 e-130
gi|928828 (L44593) ORF1904; putative [Lactococcus lactis phage B...
                                                                        427 e-118
gi|2935676 (AF032121) unknown [Streptococcus thermophilus bacter...
                                                                        309 le-82
gi|2935691 (AF032122) unknown [Streptococcus thermophilus bacter...
                                                                        306 7e-82
gi|3540289 (AF057033) putative anti-receptor [Streptococcus ther...
                                                                         279 6e-74
gi|4530154|gb|AAD21894.1| (AF085222) putative tail-host specific...
                                                                         220 3e-56
gi|930045|emb|CAA33387| (X15332) alpha-1 (III) collagen [Homo sa...
                                                                         58 4e-07
gi|1070603|pir||CGHU7L collagen alpha 1(III) chain precursor - h...
                                                                         58 4e-07
gi|4502951|ref|NP_000081.1|PCOL3A1| collagen, type III, alpha 1 ...
                                                                         58 4e-07
gi|115290|sp|P04258|CA13_BOVIN COLLAGEN ALPHA 1(III) CHAIN >gi|7...
                                                                         58 4e-07
gi|575322|emb|CAA36279| (X52046) type III collagen [Mus musculus]
                                                                         57 8e-07
gi|2119163|pir||S59856 collagen alpha 1(III) chain precursor - m...
                                                                          57 8e-07
gi|543912|sp|P13941|CA13_RAT COLLAGEN ALPHA 1(III) CHAIN >gi|543...
                                                                              1e-06
gi|3171998|emb|CAA06510| (AJ005395) collagen alpha 1 (III) [Ratt...
gi|3947565|emb|CAA90250| (Z49967) similar to collagen; cDNA EST ...
                                                                              1e-06
                                                                          57
                                                                              7e-06
                                                                          54
gi|423403|pir||A46053 bullous pemphigoid antigen, BPAG2, type XV...
                                                                          53 9e-06
gi|115410|sp|P12114|CCS1_CAEEL CUTICLE COLLAGEN SQT-1 >gi|84437|...
                                                                          53 9e-06
gi|3873801|emb|CAA90084| (Z49907) cuticle collagen SQT-1; cDNA E...
                                                                              9e-06
Query= sid|114823|lan|dp10RF002 Phage dp1 ORF|32386-35835|1
          (1149 letters)
gi|3341922|dbj|BAA31888| (AB009866) orf 15 [bacteriophage phi PVL]
                                                                         280 3e-74
gi|4126622|dbj|BAA36642.1| (AB016282) ORF36 [bacteriophage phi-105]
                                                                         232 le-59
gi|1369948|emb|CAA59194| (X84706) host interacting protein [Bact...
                                                                         201
                                                                              3e-50
gi 3139112 (AF063097) gpT [Bacteriophage P2]
gi 3337272 (U32222) G protein [Bacteriophage 186]
                                                                         188 2e-46
                                                                              3e-38
                                                                         161
gi|4063799|dbj|BAA36253| (AB008550) orf25; similar to T gene of ...
                                                                         159 8e-38
gi|3172274 (AF022214) minor tail subunit; putative tape-measure ...
                                                                         123
                                                                              6e-27
gi|465127|sp|Q05233|VG26_BPML5 MINOR TAIL PROTEIN GP26 >gi|41904...
                                                                         108
                                                                              2e-22
gi|3540284 (AF057033) putative minor tail protein (Streptococcus...
                                                                              2e-19
gi|2444119 (U88974) ORF40 (Streptococcus thermophilus temperate ...
                                                                          90
gi|2634555|emb|CAB14053| (Z99115) yomI [Bacillus subtilis] >gi|3...
                                                                          66 le-09
                                                                              5e-09
gi|2392838 (AF011378) unknown [Bacteriophage sk1]
                                                                          64
gi|2764873|emb|CAA66557| (X97918) gene 18.1 [Bacteriophage SPP1]
                                                                          62 3e-08
                                                                          61
                                                                              6e-08
gi|1353559 (U38906) ORF42 [Bacteriophage rlt]
gi|630841|pir||S39079 puff C-8 protein - fungus gnat (Rhynchosci...
                                                                              2e-06
                                                                          55
gi|1730865|sp|P51731|Y027_BPHP1 HYPOTHETICAL 72.8 KD PROTEIN IN ...
                                                                              8e-06
                                                                              1e-05
gi|224288|prf||1101273J ORF 7 [Bacteriophage HP1]
Query= sid|114824|lan|dp10RF003 Phage dp1 ORF|53538-55877|3
          (779 letters)
gi|118825|sp|P00582|DP01_ECOLI DNA POLYMERASE I (POL I) >gi|6705...
gi|2982102|pdb|1KFS|A Chain A. All-Oxygen Dna Complexed To The 3...
                                                                              3e-48
                                                                         193
                                                                              3e-48
gi|229889|pdb|1DPI| DNA Polymerase I (Klenow Fragment) (E.C.2....
                                                                         193
gi|1169402|sp|P43741|DP01_HAEIN DNA POLYMERASE I (POL I) >gi|107...
                                                                         191
                                                                              1e-47
gi|2688462 (AE001156) DNA polymerase I (polA) [Borrelia burgdorf...
                                                                         190
                                                                              3e-47
                                                                         190
                                                                              3e-47
gi|809180|pdb|1KLN|A Escherichia coli
gi|1913934|emb|CAA72997| (Y12328) DNA-directed DNA polymerase I ...
                                                                         189
                                                                               8e-47
gi|4090935 (AF028719) DNA polymerase type I (Rhodothermus sp. 'I...
                                                                         175
                                                                              1e-42
gi|4731571|gb|AAD28505.1|AF121780_1 (AF121780) DNA polymerase I ...
                                                                              2e-42
                                                                         174
gi|1633576 (U57757) similar to proofreading 3'-5' exonuclease an...
                                                                         173
                                                                              4e-42
gi|3322368 (AE001195) DNA polymerase I (polA) [Treponema pallidum]
                                                                         172
                                                                              9e-42
gi|1006595|dbj|BAA10748| (D64005) DNA polymerase I [Synechocysti...
                                                                         171
                                                                              2e-41
gi|585062|ap|Q07700|DPO1_MYCTU DNA POLYMERASE I (POL I) >gi|4161...
                                                                         163
                                                                              5e-39
gi|4376908|gb|AAD18751| (AE001645) DNA Polymerase I [Chlamydia p...
gi|1169403|sp|P46835|DP01_MYCLE DNA POLYMERASE I (POL I) >gi|107...
                                                                         157
                                                                              2e-37
                                                                         152
                                                                              7e-36
gi|2145839|pir||S72949 DNA polymerase I - Mycobacterium leprae >...
                                                                              7e-36
                                                                         152
gi|1405438|emb|CAA67184| (X98575) DNA-dependent DNA polymerase [...
                                                                        152 9e-36
gi|2506365|sp|P80194|DP01_THECA DNA POLYMERASE I, THERMOSTABLE (...
                                                                         147
                                                                              2e-34
gi|3328929 (AE001322) DNA Polymerase I [Chlamydia trachomatis]
                                                                         147 3e-34
```

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gi|3913510|sp|052225|DP01_THEFI DNA POLYMERASE I, THERMOSTABLE (...
                                                                          146 7e-34
gi|1205984 (U33536) DNA polymerase I (Bacillus stearothermophilus)
                                                                                7e-34
gi|118827|sp|P13252|DP01_STRPN DNA POLYMERASE I (POL I) >gi|9802...
                                                                           145 9e-34
                       Stoffel Fragment Of Taq Dna Polymerase I
                                                                          145 le-33
145 le-33
gi|1942202|pdb|1JXE|
gi | 1943520 | pdb | 1KTQ |
                       Dna Polymerase
gi|1084022|pir||JX0359 DNA-directed DNA polymerase (EC 2.7.7.7) ...
                                                                           145 le-33
gi|507891|dbj|BAA06775| (D32013) DNA Polymerase [Thermus aquaticus]
                                                                           145 le-33
gi | 118828 | sp | P19821 | DPO1_THEAQ DNA POLYMERASE I, THERMOSTABLE (T...
                                                                           145 le-33
gi|1706502|sp|P52028|DP01_THETH DNA POLYMERASE I, THERMOSTABLE (...
                                                                          144 2e-33
144 2e-33
gi|1097211|prf||2113329A DNA polymerase [Thermus aquaticus therm...
gi|2098289|pdb|1TAU|A Chain A, Structure Of Dna Polymerase
                                                                           143 3e-33
Query= sid|114825|lan|dp10RF004 Phage dp1 ORF|40401-42440|3
          (679 letters)
qi|1934761|emb|CAB07981| (Z93946) hypothetical protein [bacterio... 1011 0.0
gi|3540290 (AF057033) putative minor structural protein (Strepto...
gi 2444125 (U88974) ORF46 (Streptococcus thermophilus temperate ...
                                                                           339 3e-92
gi|1934762|emb|CAB07982| (Z93946) hypothetical protein [bacterio...
                                                                          300 2e-80
276 4e-73
gi|4530155|gb|AAD21895.1| (AF085222) unknown (Streptococcus ther...
gi|2935677 (AF032121) unknown (Streptococcus thermophilus bacter...
                                                                          250 3e-65
gi 2935692 (AF032122) unknown [Streptococcus thermophilus bacter...
                                                                           250 3e-65
gi 1136289 (U42597) histidine kinase A (Dictyostelium discoideum)
                                                                            50 7e-05
Query= sid|114827|lan|dp10RF006 Phage dp1 ORF|45296-46987|2
          (563 letters)
gi|4377165|gb|AAD18987| (AE001666) SWI/SNF family helicase_2 [Ch...
                                                                          171 le-41
gi|1769947|emb|CAA67095| (X98455) SNF (Bacillus cereus)
                                                                          160 3e-38
gi|3329163 (AE001341) SWF/SNF family helicase [Chlamydia trachom...
                                                                          159
                                                                               6e-38
gi|4377149|gb|AAD18973| (AE001664) SWI/SNF family helicase_1 [Ch...
                                                                          157 2e-37
                                                                          153 2e-36
gi 3328995 (AE001326) SWI/SNF family helicase [Chlamydia trachom...
gi 2493354 sp P75093 Y018 MYCPN HYPOTHETICAL HELICASE MG018/MG01...
                                                                          146 4e-34
gi|1653748|dbj|BAA18659| (D90916) helicase of the snf2/rad54 fam...
gi|1763712|emb|CAB05939| (Z83337) member of the SNF2 helicase fa...
                                                                          143 3e-33
                                                                          143 4e-33
gi|2636153|emb|CAB15645.1| (Z99122) similar to SNF2 helicase [Ba...
gi|2909552|emb|CAA17284| (AL021924) helZ [Mycobacterium tubercul...
                                                                          143 4e-33
                                                                          140 2e-32
qi 3844627 (U39681) ATP-dependent RNA helicase, putative [Mycopl...
                                                                          136 3e-31
gi | 1351463 | sp | P47264 | Y018 MYCGE HYPOTHETICAL HELICASE MG018
                                                                          136 4e-31
gi 2660669 (AC002342) human Mi-2 autoantigen-like protein [Arabi...
                                                                          131 2e-29
gi|1361537|pir||164201 helicase (mot1) homolog - Mycoplasma geni...
                                                                          129 4e-29
gi 3482977 emb CAA20533.1 (AL031369) putative protein [Arabidop...
                                                                          128 9e-29
gi|3298562 (U91543) zinc-finger helicase (Homo sapiens)
                                                                          120 2e-26
gi|3875971|emb|CAB02491| (Z80344) similar to helicase; cDNA EST ...
                                                                          120 2e-26
gi|4557451|ref|NP_001263.1|PCHD3| chromodomain helicase DNA bind...
                                                                          120 2e-26
gi|2645435 (AF007780) CHD3 [Drosophila melanogaster]
                                                                          118
                                                                               1e-25
gi 3875165 emb CAA91798 (267881) Similarity to Mouse Chromodoma...
Query= sid|114828|lan|dplORF007 Phage dp1 ORF|22230-23621|3
          (463 letters)
qi|2444105 (U88974) ORF26 [Streptococcus thermophilus temperate ...
                                                                           89 7e-17
gi|3318666 (U19754) BBA31 homolog (Borrelia burgdorferi)
                                                                                7e-08
gi|2690260 (AE000790) conserved hypothetical protein [Borrelia b...
                                                                           56 5e-07
Query= sid|114829|lan|dp10RF008 Phage dp1 ORF|49624-50961|1
gi|4406210|gb|AAD19901| (AF100420) DnaB replication fork helicas...
gi|3121983|sp|025916|DNAB_HELPY REPLICATIVE DNA HELICASE >gi|231...
                                                                               2e-10
gi|4416322|gb|AAD20314| (AF106032) replicative helicase; DnaB [B...
                                                                            65 9e-10
gi|4155895 (AE001551) REPLICATIVE DNA HELICASE (Helicobacter pyl...
                                                                            60 48-08
gi|3322317 (AE001191) replicative DNA helicase (dnaB) (Treponema...
                                                                           58 le-07
gi|138031|sp|P04530|VG41_BPT4 PRIMASE-HELICASE (PROTEIN GP41) >g...
                                                                           53 3e-06
gi|2983861 (AE000742) replicative DNA helicase (Aquifex aeolicus)
                                                                           51 1e-05
Query= sid|114831|lan|dp1ORF010 Phage dp1 ORF|8699-9859|2
          (386 letters)
                                                                                         ____
                                                                          133 2e-30 _
gi|2760912 (AF037258) RecA protein [Chlorobium tepidum]
gi 3219851 sp P94666 RECA_CLOPE RECA PROTEIN >gi 1698591 (U61497...
                                                                               3e-29
gi | 1350566 | Sp | P48295 | RECA STRVL RECA PROTEIN >gi | 508860 (U04837)...
                                                                          128
                                                                                7e-29
gi|744163|prf||2014250A recA-like protein [Streptomyces violaceus]
                                                                          126
                                                                               3e-28
gi | 730487 | sp | P41054 | RECA STRAM RECA PROTEIN >gi | 511133 | emb | CAA82...
gi | 2687334 | emb | CAA15875 | (AL020958) RecA protein [Streptomyces c...
                                                                          125
                                                                                4e-28
                                                                          125
                                                                               6e-28
gi|1350565|sp|P48294|RECA_STRLI RECA_PROTEIN >gi|481482|pir||S38...
                                                                          125 6e-28
```

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gi|464599|sp|P33542|RECA_AQUPY RECA PROTEIN >gi|1086167|pir||A55...
                                                                            123 2e-27
gi|417636|sp|P32725|RECA_RHOSH RECA_PROTEIN >gi|541307|pir||S415...
                                                                            123 2e-27
gi|2984348 (AE000775) recombination protein RecA [Aquifex aeolicus]
                                                                            123 2e-27
gi|3219854|sp|P95846|RECA_STRRM RECA PROTEIN >gi|1729800|emb|CAA...
                                                                            122 4e-27
gi|2500086|sp|Q59560|RECA_MYCSM RECA PROTEIN >gi|1430892|emb|CAA...
                                                                             122 4e-27
gi|1350567|sp|P48296|RECA_THEAQ RECA PROTEIN >gi|1072963|pir||A5...
                                                                            122 6e-27
gi|625663|pir||JX0292 recA protein - Thermus aquaticus (strain HB8)
                                                                            121 le-26
gi|1172880|sp|P42440|RECA_CAMJE RECA PROTEIN >gi|2119991|pir||14...
                                                                            120 2e-26
gi|4154654 (AE001453) RECA PROTEIN. [Helicobacter pylori J99]
                                                                             120 2e-26
gi|1072968|pir||C55020 recA protein - Thermus sp >gi|458472|dbj|...
                                                                             120 2e-26
gi|3219852|8p|P95469|RECA_PARDE RECA_PROTEIN >gi|1825468 (U59631...
                                                                            119 3e-26
gi|2507284|sp|P42445|RECA_HELPY RECA_PROTEIN >gi|2313235|gb|AAD0...
                                                                            119 4e-26
                                                                            118 5e-26
gi|1172890|sp|Q02350|RECA_STAAU RECA PROTEIN >gi|463285 (L25893)...
gi|4416209|gb|AAD20261| (AF094756) RecA protein [Bifidobacterium...
                                                                            118 5e-26
gi|2500084|Sp|Q59180|RECA_BORBU RECA PROTEIN >gi|1276443 (U23457...
                                                                            118 5e-26
Query= sid|114832|lan|dplORF011 Phage dpl ORF|28017-29096|3
          (359 letters)
gi|2444110 (U88974) ORF31 (Streptococcus thermophilus temperate ...
                                                                             187 le-46
gi|3320438 (AF057033) gp348 [Streptococcus thermophilus bacterio...
                                                                            179 2e-44
gi|479514|pir||S34244 hypothetical protein p38 - actinophage VWB...
                                                                              62 8e-09
Query= sid | 114834 | lan | dp1ORF013 Phage dp1 ORF | 10215-11240 | 3
          (341 letters)
gi|580855|emb|CAA29958| (X06803) dnaZX-like ORF put. DNA polymer...
                                                                             182 2e-45
gi|118807|sp|P09122|DP3X_BACSU DNA POLYMERASE III SUBUNITS GAMMA...
                                                                             182 2e-45
gi|98292|pir||S13786 DNA-directed DNA polymerase (EC 2.7.7.7) II...
                                                                             182 2e-45
gi|1527142 (U66040) DNA polymerase III gamma subunit [Salmonella...
gi|2494197|sp|P74876|DP3X_SALTY DNA POLYMERASE III SUBUNITS GAMM...
                                                                             172 4e-42
                                                                             172 4e-42
gi|118808|sp|P06710|DP3X_ECOLI DNA POLYMERASE III SUBUNITS GAMMA...
                                                                             170 le-41
gi|4155207 (AE001497) DNA POLYMERASE III SUBUNITS GAMMA AND TAU ...
                                                                             169 2e-41
gi|2313841|gb|AAD07767.1| (AE000584) DNA polymerase III gamma an...
                                                                             168 4e-41
gi|2583049 (AF025391) DNA polymerase III holoenzyme tau subunit ...
gi|2984127 (AE000759) DNA polymerase III gamma subunit [Aquifex ...
                                                                             166 3e-40
                                                                             166 3e-40
165 5e-40
gi 3861390 emb CAA15289 (AJ235273) DNA POLYMERASE III SUBUNITS ... gi 1169397 sp P43746 DP3X HAEIN DNA POLYMERASE III SUBUNITS GAMM...
                                                                             156 2e-37
gi|1293572 (U49738) DNA polymerase III tau homolog DnaX [Cauloba...
                                                                             151 8e-36
gi|3328753 (AE001306) DNA Pol III Gamma and Tau (Chlamydia trach...
gi|4376294|gb|AAD18193| (AE001589) DNA Polymerase III Gamma and ...
gi|581255|emb|CAA28175| (X04487) alternate dnaZX protein (AA 1-6...
                                                                             148
                                                                                  5e-35
                                                                             146 3e-34
gi 2688379 (AE001151) DNA polymerase III, subunits gamma and tau...
                                                                             140 2e-32
gi|3323329 (AE001268) DNA polymerase III, subunits gamma and tau...
                                                                             137 le-31
Query= sid|114835|lan|dp10RF014 Phage dp1 0RF|50961-51974|3
          (337 letters)
gi|1346796|sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64...
                                                                              57 2e-07
gi|740008|prf||2004290A primase [Haemophilus influenzae]
                                                                              51 le-05
gi|1172619|sp|Q08346|PRIM_HABIN DNA PRIMASE >gi|1074033|pir||A64...
                                                                              51 le-05
gi | 1709769 | sp | Q04505 | PRIM_LACLA DNA PRIMASE >gi | 1075726 | pir | | JC2...
                                                                              51 1e-05
                                                                              51 le-05
gi|639846|dbj|BAA03516| (D14690) DNA primase [Lactococcus lactis]
Query= sid|114837|lan|dp1ORF016 Phage dp1 ORF|43413-44303|3
          (296 letters)
gi|1934766|emb|CAB07986| (293946) N-acetylmuramoyl-L-alanine ami...
                                                                             661 0.0
gi 113676 sp | P06653 | ALYS_STRPN AUTOLYSIN (N-ACETYLMURAMOYL-L-ALA...
                                                                             221
gi|282326|pir||A42935 N-acetylmuramoyl-L-alanine amidase (EC 3.5...
                                                                             219 3e-56
                                                                             212 2e-54
212 2e-54
gi 416618 sp | P32762 | ALYS_BPHB3 LYTIC AMIDASE (N-ACETYLMURAMOYL-L...
gi|285273|pir||A42936 N-acetylmuramoyl-L-alanine amidase (EC 3.5...
gi|127787|sp|P15057|LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE)...
                                                                             162 4e-39
gi|67761|pir||MUBPCP N-acetylmuramoyl-L-alanine amidase (EC 3.5....
                                                                             162 4e-39
gi|127789|sp|P19386|LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE)...
gi|928832 (L44593) ORF259; putative (Lactococcus lactis phage BK...
                                                                             119 2e-26
gi|2511705|emb|CAA71783| (Y10818) sigA binding protein (Streptoc...
                                                                             111 9e-24
                                                                             107 le-22
105 4e-22
gi 4097980 (U72655) surface protein C [Streptococcus pneumoniae]
gi|2351768 (U89711) PspA (Streptococcus pneumoniae)
                                                                                            ____
gi|2425109 (AF019904) choline binding protein A [Streptococcus p...
                                                                             104 6e-22
                                                                             104 1e-21
103 2e-21
gi|282335|pir||A41971 surface protein pspA precursor - Streptoco...
gi|2576331|emb|CAA05158| (AJ002054) SpsA protein (Streptococcus ...
                                                                              85 6e-16
84 1e-15
gi|2127295|pir||S57962 cspC protein - Clostridium acetobutylicum...
gi|2576333|emb|CAA05159| (AJ002055) SpsA protein (Streptococcus ...
gi|4106522|gb|AAD02874.1| (AF097909) excreted protein FibB [Pept...
gi|1361406|pir||557714 cspB protein - Clostridium acetobutylicum...
gi|1914872|emb|CAB04758| (Z82001) PCPA [Streptococcus pneumoniae]
                                                                              83 3e-15
82 4e-15
                                                                              81 9e-15
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gi|3168594|dbj|BAA28613| (AB012763) SpaA (Erysipelothrix rhusiop...
                                                                                      81 le-14
80 3e-14
gi|2292750|emb|CAA64942| (X95646) homology to orf259 of lactococ...
gi|2935696 (AF032122) putative lysin [Streptococcus thermophilus...
                                                                                      80 3e-14
80 3e-14
79 5e-14
gi|4586910|dbj|BAA76540.1| (AB017447) protective antigen SpaA.1 ...
gi|3540294 (AF057033) lysin (Streptococcus thermophilus bacterio...
Query= sid|114841|lan|dp1ORF020 Phage dp1 ORF|1864-2658|1
           (264 letters)
gi|2633745|emb|CAB13247| (Z99111) similar to coenzyme PQQ synthe...
gi|2808502|emb|CAA12532| (AJ225561) ExsD protein [Sinorhizobium ...
gi|3861151|emb|CAA15051| (AJ235272) unknown [Rickettsia prowazekii]
gi|1652793|db||BAA17712| (D90908) hypothetical protein [Synechoc...
                                                                                     163 le-39
82 6e-15
                                                                                       76 3e-13
70 2e-11
gi|1723815|sp|P55139|YGCF_ECOLI HYPOTHETICAL 25.0 KD PROTEIN IN ...
gi 2984272 (AE000769) hypothetical protein [Aquifex aeolicus]
gi 4155435 (AE001516) putative [Helicobacter pylori J99]
                                                                                       66 4e-10
                                                                                       57 1e-07
gi|2127833|pir||C64505 coenzyme PQQ synthesis protein III homolo...
                                                                                       55 Se-07
gi|2622338 (AE000890) coenzyme PQQ synthesis protein III [Methan...
                                                                                       54 9e-07
53 2e-06
gi|3257042|dbj|BAA29725| (AP000003) 254aa long hypothetical prot...
gi|2314068|gb|AAD07976.1| (AE000602) conserved hypothetical prot...
gi|1723816|sp|P45097|YGCF_HAEIN HYPOTHETICAL PROTEIN HI1189 >gi|...
                                                                                      52 6e-06
                                                                                     50 2e-05
Query= sid|114842|lan|dp10RF021 Phage dp1 0RF|2504-3295|2
            (263 letters)
gi|127481|sp|P19465|GCH1_BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >...
                                                                                      208 4e-53
gi 3242315 emb CAA04237 (AJ000685) GTP cyclohydrolase (Streptoc...
                                                                                      191 4e-48
gi|2494695|sp|Q54769|GCH1_SYNP7 GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                            2e-47
                                                                                      189
gi 255061|bbs 112832 (S44049) GTP cyclohydrolase I (clone hGCH-1...
gi 4503949 ref NP 000152.1|PGCH1| GTP cyclohydrolase 1 (dopa-res...
                                                                                      187
                                                                                            7e-47
                                                                                      187 7e-47
gi|2113967|emb|CAB08935| (Z95557) folE [Mycobacterium tuberculosis]
                                                                                      187
gi 1730240 sp P50141 GCH1 CHICK GTP CYCLOHYDROLASE I (GTP-CH-I) ...
gi 2494696 sp Q55759 GCH1_SYNY3 GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                      185 3e-46
                                                                                      184 5e-46
gi|121061|sp|P22288|GCH1_RAT GTP CYCLOHYDROLASE I PRECURSOR (GTP...
                                                                                      184 6e-46
 gi|3183014|sp|013774|GCHI_SCHPO GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                      184 6e-46
gi 3097224 emb | CAA18795 | (AL023093) GTP cyclohydrolase I (Mycoba...
                                                                                      182 2e-45
gi 2494697 Bp |Q19980 |GCH1 CAEEL PROBABLE GTP CYCLOHYDROLASE I (G... gi 462167 |Bp |Q05915 |GCH1 MOUSE GTP CYCLOHYDROLASE I PRECURSOR (G...
                                                                                      182 2e-45
                                                                                      180 7e-45
180 1e-44
gi|1669664|emb|CAA89808| (Z49706) GTP cyclohydrolase I [Dictyost...
                                                                                      178 3e-44
177 8e-44
gi|2981082 (AF052048) GTP-cyclohydrolase [Ostertagia ostertagi]
gi|31954|emb|CAA78908| (Z16418) GTP cyclohydrolase I [Homo sapi...
 gi|551344|bbs|150280 (S71373) GTP cyclohydrolase I [mice, Peptid...
                                                                                     174 5e-43
174 7e-43
 gi | 1730247 | sp | P51601 | GCH1_YEAST GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                     172 2e-42
 gi | 1246912 emb | CAA87397 | (Z47201) GTP cyclohydrolase 1 [Saccharo...
 gi 1730246 sp P51595 GCH1 STRPN GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                     168 3e-41
 gi|2982951 (AE000680) GTP cyclohydrolase I [Aquifex aeolicus]
                                                                                      164 6e-40
 Query= sid|114843|lan|dp10RF022 Phage dp1 ORF|30896-31675|2
            (259 letters)
 gi|2347102 (U77367) internalin [Listeria monocytogenes]
gi|3123226|sp|P25146|INLA_LISMO INTERNALIN A PRECURSOR >gi|48705...
                                                                                        55 5e-07
                                                                                       52 4e-06
                                                                                        52 4A-06
 gi|149674 (M67471) internalin [Listeria monocytogenes]
 Query= sid|114850|lan|dp1ORF029 Phage dp1 ORF|662-1348|2
            (228 letters)
 gi|2650185 (AE001074) succinoglycan biosynthesis regulator (exsB...
                                                                                      119 2e-26
 gi|3861231|emb|CAA15131| (AJ235272) unknown [Rickettsia prowazekii]
                                                                                      117 8e-26
 gi|2622210 (AE000881) conserved protein [Methanobacterium thermo...
                                                                                      108 4e-23
 gi|2983380 (AE000709) trans-regulatory protein ExsB (Aquifex aeo...
                                                                                      88 6e-17
 gi|1001327|dbj|BAA10814| (D64006) ExsB [Synechocystis sp.]
 gi|2128055|pir||B64468 hypothetical protein homolog MJ1347 - Met...
                                                                                       83 le-15
                                                                                       82 4e-15
80 2e-14
 gi 4155143 (AE001491) putative [Helicobacter pylori J99]
 gi|2313760|gb|AAD07701.1| (AE000578) conserved hypothetical prot...
gi|2120814|pir||S60183 protein ExsB - Rhizobium meliloti >gi|114...
                                                                                       76 3e-13
 gi|2633743|emb|CAB13245| (Z99111) similar to hypothetical protei...
                                                                                       75 5e-13
 gi 1175543 sp P44124 YBAX HAEIN HYPOTHETICAL PROTEIN HI1191 >gi ...
gi 2495537 sp P77756 YBAX ECOLI HYPOTHETICAL 25.5 KD PROTEIN IN ...
                                                                                      74 1e-12
                                                                                     71 5e-12 --- --- 67 1e-10 ---
 gi|3256471|dbj|BAA29154.1| (AP000001) 269aa long hypothetical pr...
                                                                                     54 le-06
 gi|2921156 (AF022216) aluminum resistance protein (Arthrobacter ...
 Query= sid|114855|lan|dp10RF034 Phage dp1 ORF|131-652|2
            (173 letters)
 gi|2633746|emb|CAB13248| (Z99111) similar to hypothetical protei... 220 4e-57
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717		
gi[4155926 (AE001554) putative [Helicobacter pylori J99]	162	1e-39
gi 2314588 gb AAD08456.1  (AE000642) conserved hypothetical prot	161	3e-39
gi 2983458 (AE000714) hypothetical protein [Aquifex aeolicus]	103	9e-22
gi 1006604 dbj BAA10757  (D64005) hypothetical protein (Synechoc	87	6e-17
gi 2967529 (Ul1045) unknown (Buchnera aphidicola)	79	2e-14
gi 2495654 sp Q46920 YQCD_ECOLI HYPOTHETICAL 32.6 KD PROTEIN IN	69	2e-11
g1 2495654 BP Q46920 YCCD ECOLI HIPOTHETICAL 32.6 KD PROTEIN IN		
gi 1175604 sp P44153 YQCD HAEIN HYPOTHETICAL PROTEIN HI1291 >gi	63	1e-09
gi 3860642 emb CAA14543  (AJ235270) unknown [Rickettsia prowazekii]	56	1e-07
Query= sid 114857 lan dp10RF036 Phage dp1 ORF 48808-49362 1 (184 letters)		
gi 1353529 (U38906) ORF12 [Bacteriophage rlt]	53	1e-06
Query= sid 114859 lan dp10RF038 Phage dp1 ORF 1350-1871 3 (173 letters)		
gi 1175542 sp P44123 YB90_HAEIN HYPOTHETICAL PROTEIN HIll90 >gi	100	6e-21
gi 2982977 (AE000681) hypothetical protein (Aquifex aeolicus)	67	7e-11
gi 3860744 emb CAA14645  (AJ235270) unknown [Rickettsia prowazekii]	65	3e-10
gi 2650193 (AE001074) conserved hypothetical protein (Archaeoglo	58	4e-08
gi 3258383 dbj BAA31066.1  (APO00007) 157aa long hypothetical pr	55	2e-07
g1 3258383 GD BAA31066.1 (APOUDOV) 15/44 Tong hypothetical pr	50	8e-06
gi 1001713 dbj BAA10550 (D64004) hypothetical protein [Synechoc		
gi 4155434 (AE001516) putative (Helicobacter pylori J99)	50	1e-05
Query= sid 114860 lan dp10RF039 Phage dp1 ORF 3306-3803 3 (165 letters)		
gi 1922884 emb CAA68244  (X99978) ORF7; hydophobic protein [Lact	64	5e-10
Query= sid 114862 lan dp10RF041 Phage dp1 ORF 8208-8699 3 (163 letters)		
gi 2522313 (AF012906) dUTPase homolog [Bacillus subtilis] >gi 26	108	2e-23
gi 2634150 emb CAB13650 (299113) similar to deoxyuridine 5'-tri	108	3e-23
gi 3913546 sp   054134   DUT_STRCO DEOXYURIDINE 5'-TRIPHOSPHATE NUCL	56	2e-07
gi 3913542 sp 048500 DUT_BPTS DEOXYURIDINE 5'-TRIPHOSPHATE NUCLE	52	3e-06
gi 3913548 sp 068992 DUT_CHLTE DEOXYURIDINE 5'-TRIPHOSPHATE NUCL	50	1e-05
91/391346/88/000992/801_0 2201.01.2	•	
Query= sid 114867 lan dp10RF046 Phage dp1 ORF 42774-43202 3 (142 letters)		
gi 1934764 emb CAB07984  (Z93946) hypothetical protein [bacterio	287	2e-77
Query= sid 114901 lan dplORF080 Phage dpl ORF 42490-42759 1 (89 letters)		
gi 1934763 emb CAB07983  (Z93946) hypothetical protein [bacterio	147	1e-35
Query= sid 114912 lan dp10RF091 Phage dp1 ORF 43189-43413 1 (74 letters)		
gi 1934765 emb CAB07985  (Z93946) holin [bacteriophage Dp-1]	63	2e-10

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Table 32

# Sequence of Dp1 published by Sheehan and al.. 4731 nucleotides.

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141	agaccttaaa	tatcgaattg	actcaaaagc	cgatcaaaag	ctaactaacc	aacagttgac	ggcactcacg	
211	gaaaaggctc	aactacatga	cgcagaactg	aaagctaagg	ctacaatgga	gcagttaagt	aacttagaaa	
281	aggcttatga	aggtagaatg	aaagctaatg	aagaagctat	caacaaatcg	gaacccgacc	taatcttagc	
351	ggcaagtcga	attgaagcta	ctatccaaga	acttggcggg	ctacgggaac	tgaagaagtt	cgtcgacagt	
421	tgcatgagct	cttctaatca	aggtctaatt	atcggtaaga	acgacggtag	ctctaccatt	aaggtatcaa	
491	gtgaccgaat	ttctatgttc	tccgcaggga	atgaagttat	gtacettacg	caagggttca	ttcacatcga	
561	taacgggatc	tttacccaat	ccattcaagt	cggccgattt	agaacggaac	aatactcgtt	caacccagac	
631	atgaacgtga	ttcggtatgt	aggataagga	gaataacatg	acadaactta	tcaactcata	netteeeee	
701	cacttgaacc	tttacgtcga	acaagttagt	caggacgtaa	togaacaactc	ctcgcgagtt	ccatatoatt	
771	ctactgtcga	ccgcgatgga	gettategaa	egeggaeera	aceteceee	agtaaccttt aagaggtaac	acteacaget	
841	aaatggttca	agegeeata	geagceaece	agactacgac	castetees	ttgggcttcg	tttcaccta	
911	ggagaagtga	tenegateetea	arcactatct	ctactaatta	cactttagac	agtattccaa	ogtotacaca	
981	ataacggcgt	tttaaggaaat	atcactacct	accetcatta	catacogtta	tctttaaccg	aaaagtgaac	
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1611	gaaaacaatc	gaacgtccaa	gacgtatcta	tcaatgttat	agaatactat	ggaccgtcta	tcaatttctc	
1681	cottcaacqt	actcqtcaaa	atcctgcaat	tatccaagct	cttcgaaatg	ctaaggtcgc	acctataacg	
1751	gtaggaggtc	aacagaaaaa	catcatgcaa	attaccttct	ccgtggcgcc	gttgaacact	actaatttca	
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2171	agcgtgaaac	agagtttaca	tggcgaagta	acaaatacga	ggacaaccct	acgggaactc	gaggtgaatg	
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2451	cgacgcattc	tattcgaaaa	ctcttgacgg	catagtatat	ttgagaggaa	atgtgcataa	aggacttate	
2521	gacaaagagg	ctactattgc	agtacttcct	gaaggattta	gaccgaaagt	ttcaatgtat	antoneetot	
2591	tcaataactc	atatggaaat	gecattetat	geacacacac	cyacyyaaya	cttgtggtga	atottataat	
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3291	taccotcttt	atcacqactt	aaaaagggaa	gtgataacag	gctatacaac	tctcgaccat	tttagagagc	
3361	tetetatttt	attcqaaaqt	tataagaacc	ttggcggaaa	tggtgaagtt	gaageettgt	atgaaaaata	
3431	caaqaaatta	ccaattaggg	aggaagattt	agatgaaact	atctaacgaa	caatatgacg	tagcaaagaa	
3501	cqtqqtaacc	gtagtcgttc	cagcagcgat	tgcactaatt	acaggtettg	gagcgttgta	tcaatttgac	
3571	actactgcta	tcacaggaac	cattgcactt	cttgcaactt	ttgcaggtac	tgttctagga	gtttctagcc	
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#### Table 33

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gi|295191|gb|L15190.1|STRPURISYN [295191]

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### **CLAIMS**

### What is claimed is:

1. A method for identifying a bacteriophage coding region encoding a product active on an essential bacterial target, comprising identifying a nucleic acid sequence encoding a gene product which provides a bacteria-inhibiting function when said bacteriophage infects a host bacterium,

wherein said bacteriophage is uncharacterized and said host bacterium is a pathogenic bacterium.

- 2. The method of claim 1, further comprising expressing a recombinant bacteriophage ORF in cells of a bacterial strain, wherein inhibition of said cells following expression of said ORF is indicative that said product is active on an essential bacterial target.
- 3. The method of claim 2, wherein inhibition of said bacterium following expression of said ORF is determined by comparison with the growth or viability of said bacterium following expression of an inactivated mutant form of said ORF or in the absence of expression of said ORF, and wherein inhibition of said bacterium following expression of said ORF is indicative that said product is active on an essential bacterial target.
  - 4. The method of claim 2, wherein expression of said ORF is inducible.

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- 5. The method of claim 1, further comprising sequencing at least a portion of a bacteriophage genome.
- 6. The method of claim 1, wherein at least a portion of the nucleotide sequence of a bacteriophage genome is known, said method further comprising identifying at least one ORF in said portion by computer analysis of said sequence.
- 7. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify
  homologous genes or gene products of known biochemical function, therebyindicating the biochemical function of said polypeptide.

- 8. The method of claim 7, wherein said homologous gene or gene product is a bacterial gene important for cell viability.
- 9. The method of claim 7, wherein said homologous gene or gene product is a gene or gene product known to have a bacteria-inhibiting function.
  - 10. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify structural motifs in said polypeptide, thereby indicating the cellular function of said polypeptide.

- 11. The method of claim 1, wherein a host bacterium for said bacteriophage is selected from the species group consisting of bacteria listed in Table 1.
- 15 12. The method of claim 1, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.
  - 13. The method of claim 2, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.

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- 14. The method of claim 13, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.
- The method of claim 14, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.
  - 16. The method of claim 1, wherein said pathogenic bacterium is an animal pathogen.
- The method of claim 16, wherein said pathogenic bacterium is a human pathogen.
  - 18. The method of claim 1, wherein said pathogenic bacterium is a plant pathogen.

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19. The method of claim 1, further comprising confirming the inhibitor function of said ORF.

- 20. The method of claim 19, wherein said confirming comprises expressing a loss-of-function mutant form of said ORF in said host bacterium.
- 5 21. The method of claim 1, wherein said identifying a nucleic acid sequence encoding a gene product active on an essential bacterial target comprises identifying a nucleic acid sequence encoding a homolog of a bacteriophage polypeptide known to be active on an essential bacterial target.
- 10 22. The method of claim 1, wherein said identifying a bacteriophage coding region comprises identifying a first coding region from a bacteriophage having a non-pathogenic host bacterial strain related to said pathogenic bacterium, said first coding region encoding a product active on an essential bacterial target; and identifying a homolog of said first coding region, wherein said homolog is a probable said bacteriophage coding region encoding a product active on an essential bacterial target.
  - 23. The method of claim 2, wherein a plurality of bacteriophage ORFs from a plurality of different bacteriophage are expressed in at least one bacterium.
  - 24. The method of claim 23, wherein each of said plurality of bacteriophage ORFs are expressed in different bacteria.
- 25. A method for identifying a target for antibacterial agents, comprising determining the bacterial target of an uncharacterized bacteriophage inhibitor protein.
- 26. The method of claim 25, wherein said determining comprises identifying at least one bacterial protein which binds to said bacteriophage inhibitor protein or a fragment thereof.
  - 27. The method of claim 26, wherein said binding is determined using affinity chromatography on a solid matrix.
- The method of claim 25, wherein said determining comprises identifying at least one protein:protein interaction using a genetic screen.

- 29. The method of claim 28, wherein said genetic screen is a yeast twohybrid screen.
- 30. The method of claim 25, wherein said determining comprises a coimmunoprecipitation assay or a protein-protein crosslinking assay.
  - 31. The method of claim 25, wherein said determining comprises identifying a mutated bacterial coding sequence which protects a bacterium from said bacteriophage inhibitor.

- 32. The method of claim 25, wherein said determining comprises identifying a bacterial coding sequence which protects a bacterium against said bacteriophage inhibitor when expressed at high levels in said bacterium.
- 15 33. The method of claim 25, wherein said determining further comprises identifying a bacterial nucleic acid sequence encoding a polypeptide target of said bacteriophage inhibitor protein.
- 34. The method of claim 33, wherein said nucleic acid sequence is
  identified by determining at least a portion of the amino acid sequence of a bacterial protein target, and identifying a bacterial nucleic acid sequence which encodes said protein target.
- The method of claim 25, wherein said bacterial target is naturally produced by a bacterial species selected from the group consisting of species of the genera listed in Table 1.
  - 36. The method of claim 25, wherein said bacterial target is naturally produced by a bacterial strain selected from the group consisting of species listed in Table 1.
  - 37. The method of claim 25, wherein said inhibitor protein is naturally produced by a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

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38. The method of claim 25, further comprising identifying a bacteriophage ORF which encodes a product having a bacteria-inhibiting function.

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- 39. The method of claim 38, wherein said identifying a phage ORF comprises expressing at least one bacteriophage ORF in a bacterium, wherein inhibition of said bacterium following said expression is indicative that said ORF encodes a bacteria-inhibiting function.
- 40. The method of claim 39, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.
- 10 41. The method of claim 40, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.
  - 42. The method of claim 41, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.
  - 43. The method of claim 25, wherein said determining the bacterial target of a bacteriophage inhibitor protein is performed for a plurality of different bacteriophage of the same host bacterium.
- 20 44. The method of claim 25, wherein said bacterial target originates from an animal pathogen.
  - 45. The method of claim 44, wherein said bacterial target is a gene homologous to a gene from an animal pathogen.
    - 46. The method of claim 44, wherein said pathogen is a human pathogen.
  - 47. The method of claim 25, wherein said bacterial target originates from a plant pathogen.
  - 48. The method of claim 25, wherein said bacterial target is a gene homologous to a gene from a plant pathogen.
- 49. The method of claim 25, further comprising determining the cellular or \_\_\_\_\_ biochemical function or both of said inhibitor protein.

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- 50. The method of claim 25, wherein said identifying the bacterial target comprises identifying a phage-specific site of action.
- 5 51. An isolated, purified, or enriched nucleic acid sequence at least 15 nucleotides in length, wherein said sequence corresponds to at least a portion of a bacteriophage sequence, and wherein said bacteriophage is selected from the group consisting of Staphylococcus aureus bacteriophage 77, 3A, 96, and 44AHJD, Enterococcus baceriophage 182, and Streptococcus pheumoniae bacteriophage Dp-1.

52. The nucleic acid sequence of claim 51, wherein said sequence comprises at least 50 nucleotides.

- 53. The nucleic acid sequence of claim 51, wherein said nucleic acid sequence corresponds to at least a portion of a nucleic acid sequence which encodes a product which provides a bacteria-inhibiting function.
  - 54. The nucleic acid sequence of claim 53, wherein said nucleic acid sequence encodes a polypeptide which provides a bacteria-inhibiting function.
  - 55. The nucleic acid sequence of claim 54, wherein said nucleic acid sequence is transcriptionally linked with regulatory sequences enabling induction of expression of said sequence.
  - 56. An isolated, purified, or enriched polypeptide comprising at least a portion of a protein providing a bacteria-inhibiting function, wherein said polypeptide is normally encoded by a bacteriophage selected from the group consisting of Staphylococcus aureus bacteriophage 77, 3A, 96, and 44AHJD, Enterococcus baceriophage 182, and Streptococcus pheumoniae bacteriophage Dp-1.
  - 57. The polypeptide of claim 56, wherein said polypeptide provides said bacteria-inhibiting function.
- 35 58. The polypeptide of claim 56, wherein said polypeptide comprises a portion at least 10 amino acid residues in length of a said polypeptide normally encoded by said bacteriophage.

- 59. A recombinant vector comprising a bacteriophage ORF corresponding to an ORF from a bacteriophage having a pathogenic bacterial host, wherein said
   5 bacterial host is selected from the group consisting of uncharacterized bacteria of Table 1.
  - 60. The vector of claim 59, wherein said vector is an expression vector.
- 10 61. The vector of claim 59, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage of Table 1.
  - 62. The vector of claim 61, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* baceriophage 182, and *Streptococcus pheumoniae* bacteriophage Dp-1.
    - 63. The vector of claim 60, wherein expression of said ORF is inducible.
- 20 64. A recombinant cell comprising a vector, wherein said vector comprises an ORF from a bacteriophage having a pathogenic bacterial host, wherein said bacterial host is selected from the group consisting of bacterial species of Table 1.
- 65. The recombinant cell of claim 64, wherein said bacteriophage is selected from the group consisting of uncharacterized phage of Table 1.
  - 66. The cell of claim 65, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* baceriophage 182, and *Streptococcus pheumoniae* bacteriophage Dp-1.
  - 67. The cell of claim 64, wherein said vector is an expression vector and expression of said ORF is inducible.
- 35 68. A method for identifying an antibacterial agent, comprising identifying an active portion of a product of a bacteria-inhibiting ORF of a bacteriophage.

69. The method of claim 68, further comprising constructing a synthetic peptidomimetic molecule, wherein the structure of said molecule corresponds to the structure of said active portion.

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70. A method for identifying a compound active on a target of a bacteriophage inhibitor protein, comprising the step of

contacting a bacterial target protein with a test compound; and
determining whether said compound binds to or reduces the level of
activity of said target protein,

wherein binding of said compound with said target protein or a reduction of the level of activity of said protein is indicative that said compound is active on said target and wherein said target is uncharacterized.

- 71. The method of claim 70, wherein said contacting is carried out in vitro.
- 72. The method of claim 70, wherein said contacting is carried out *in vivo* in a cell.
- The method of claim 70, wherein said compound is a small molecule.
  - 74. The method of claim 70, wherein said compound is a peptidomimetic compound.
- 25 75. The method of claim 70, wherein said compound is a fragment of a bacteriophage inhibitor protein.
  - 76. The method of claim 70, further comprising determining the site of action of said compound on said target protein.

- 77. The method of claim 70, wherein said contacting is performed for a plurality of said target proteins.
- 35 78. A method of screening for potential antibacterial agents, comprising the step of determining whether any of a plurality of compounds is active on a target of a bacteriophage inhibitor protein,

wherein said target is naturally produced by a pathogenic bacterium.

- 79. The method of claim 78, wherein said plurality of compounds are small molecules.
- 80. The method of claim 78, wherein said determining is performed for a plurality of said targets.
- 10 81. A method for inhibiting a bacterium, comprising the step of; contacting said bacterium with a compound active on a target of a bacteriophage inhibitor protein, wherein said target or the target site is uncharacterized.
- 15 82. The method of claim 81, wherein said compound is said protein or an active fragment thereof.
  - 83. The method of claim 81, wherein said compound is a structural mimetic of said protein.
    - 84. The method of claim 81, wherein said compound is a small molecule.
    - 85. The method of claim 81, wherein said contacting is performed in vitro.
- 25 86. The method of claim 81, wherein said contacting is performed *in vivo* in an animal.
  - 87. The method of claim 86, wherein said animal is a human.
- 30 88. The method of claim 81, wherein said contacting is carried out *in vivo* in a plant.
  - 89. The method of claim 81, wherein said bacterium is selected from the group of bacteria listed in Table 1.

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- 90. A method for treating a bacterial infection in an animal suffering from an infection, comprising administering to said animal a therapeutically effective amount of compound active on a target of a bacteriophage inhibitor protein in a bacterium involved in said infection,
- wherein said target is an uncharacterized target or the compound is active at an uncharacterized target site.
  - 91. The method of claim 90, wherein said compound is a small molecule.
- 10 92. The method of claim 90, wherein said compound is a peptidomimetic compound.
  - 93. The method of claim 90, wherein said compound is a fragment of a bacteriophage inhibitor protein.
    - 94. The method of claim 90, wherein said animal is a mammal.
    - 95. The method of claim 94, wherein said mammal is a human.
- 20 96. The method of claim 90, wherein said bacterium is selected from the group listed in Table 1.
  - 97. The method of claim 90, wherein said bacteriophage inhibitor protein is from a bacteriophage selected from the group of bacteriophage listed in Table 1.
  - 98. A method for propylactically treating an animal at risk of an infection, comprising administering to said animal a prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein,
- wherein said target is an uncharacterized target or the site of action of said compound is an uncharacterized target site.
  - 99. The method of claim 98, wherein said compound is a small molecule.
- 35 100. The method of claim 98, wherein said compound is a peptidomimetic compound.

- 101. The method of claim 98, wherein said compound is a fragment of a bacteriophage inhibitor protein.
  - 102. The method of claim 98, wherein said animal is a mammal.

- 103. The method of claim 102, wherein said mammal is a human.
- 104. An antibacterial agent active on a target of a bacteriophage inhibitor protein, wherein said target is an uncharacterized target or said agent is active at a phage-specific site on said target.
  - 105. The agent of claim 104, wherein said agent is a pepetidomimetic of a bacteriophage inhibitor polypeptide.

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- 106. The agent of claim 104, wherein said agent is a small molecule.
- 107. The agent of claim 104, wherein said agent is a fragment of a bacteriophage inhibitor polypeptide.

- 108. The agent of claim 104, wherein said agent is active at a phage-specific site on said target.
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- 109. A method of making an antibacterial agent, comprising the steps of:
  - a) identifying a target of a bacteriophage inhibitor polypeptide;
- b) screening a plurality of test compounds to identify a compound active on said target; and
- c) synthesizing said compound in an amount sufficient to provide a therapeutic effect when administered to an organism infected by a bacterium naturally producing said target.
  - 110. The method of claim 109, wherein said compound is a small molecule.
- The method of claim 109, wherein said compound is a peptidemimetic compound.

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- 112. The method of claim 109, wherein said compound is a fragment or derivative of a bacteriophage inhibitor protein.
- 113. A computer readable device having recorded therein a nucleotide sequence of a portion of at least one bacteriophage genome of *Staphylococcus aureus* bacteriophage 77, bacteriophage 3A, or bacteriophage 96, a nucleotide sequence at least 95% identical to a said nucleotide sequence, a ribonucleic acid equivalent, a degenerate equivalent, a homologous sequence, or at least one amino acid sequence encoded by said nucleotide sequence; and

a nucleotide sequence or amino acid sequence analysis program,
wherein said program can perform at least one sequence analysis on said
nucleotide or amino acid sequence.

- 15 114. The device of claim 113, wherein said at least a portion of at least one bacteriophage genome comprises at least one ORF.
  - 115. The device of claim 113, wherein said device comprises a medium selected from the group consisting of floppy disk, computer hard drive, optical disk, computer random access memory, and magnetic tape wherein said nucleotide or amino acid sequence or said program or both are recorded on said medium.
  - 116. The device of claim 113, wherein said portion of at least one bacteriophage genomic nucleotide sequence comprises at least 50% of at least one bacteriophage genomic sequence.
  - 117. The device of claim 113, wherein said at least one bacteriophage nucleotide genomic sequence comprises portions of a plurality of bacteriophage nucleotide genomic sequences.

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- 118. A computer-based system for identifying biologically important portions of a bacteriophage genome, comprising:
- a) a data storage medium having recorded thereon a nucleotide sequence
   corresponding to a portion of at least one bacteriophage genome, wherein said
   bacteriophage genome is uncharacterized;

- b) a set of instructions allowing searching of said sequence to analyze said sequence; and
  - c) an output device.
- 5 119. The system of claim 118, wherein said output device comprises comprises a device selected from the group consisting of a printer, a video display, and a recording medium.
- 120. The system of claim 118, wherein said bacteriophage genome is of a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.
  - 121. The system of claim 118, wherein said uncharacterized bacteriophage is selected from the group consisting of bacteriophage 77, 3A, and 96.

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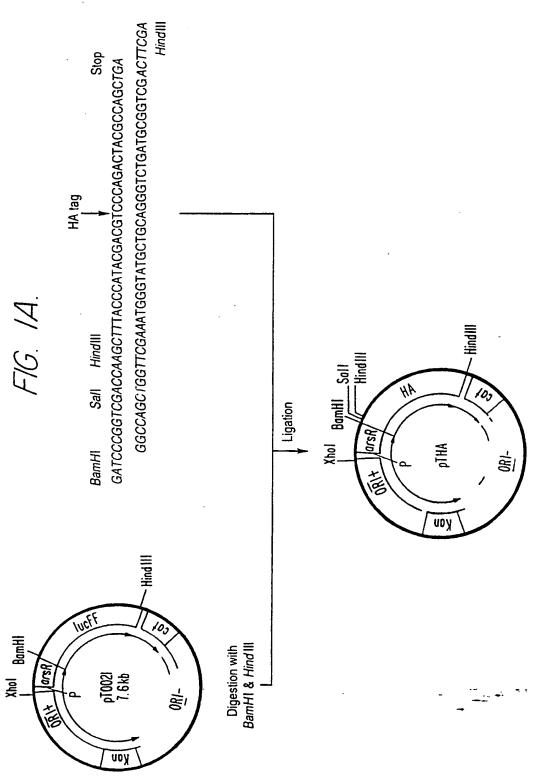
- 122. A method for identifying or characterizing a bacteriophage ORF, comprising the steps of:
- a) providing a computer-based system for analyzing nucleic acid or
  20 amino acid sequence data, wherein said system comprises a data storage medium
  having recorded thereon at least one nucleotide or amino acid sequence corresponding
  to a portion of at least one uncharacterized bacteriophage genome, a set of instructions
  allowing searching of said sequence to analyze said sequence; and an output device;
  - b) analyzing at least a portion of at least one said sequence; and
  - c) outputting results of said analyzing to said output device.
  - 123. The method of claim 122, wherein said analysis identifies sequence similarity or homology with sequences selected from the group consisting of bacterial ORFs encoding products with related biological function; ORFs encoding known inhibitors or bacteria, essential bacterial ORFs.
  - 124. The method of claim 122, wherein said analysis comprises identifying a probable biological function based on identification of structural elements or sequence homology or similarity.

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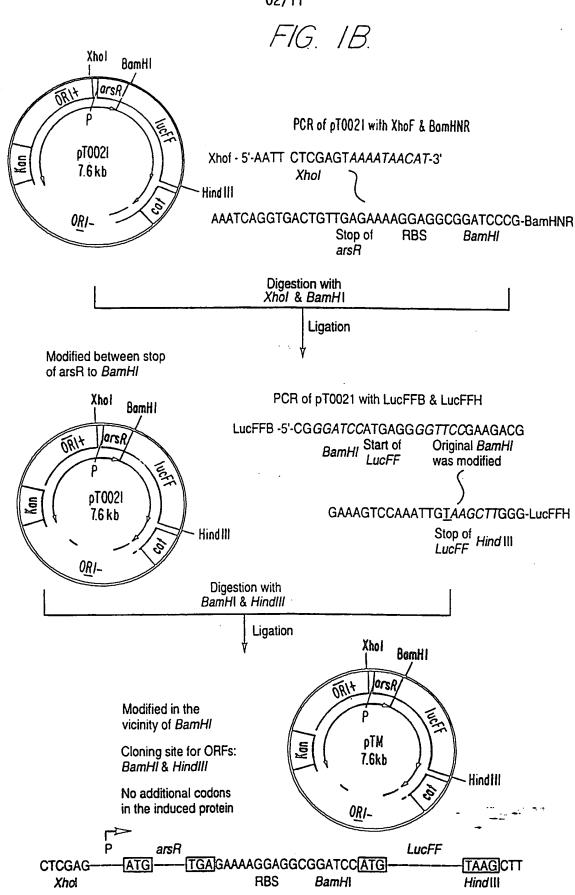
125. The method of claim 122, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

126. The method of claim 125, wherein said uncharacterized bacteriophage is selected from bacteriophage 77, 3A, and 96.



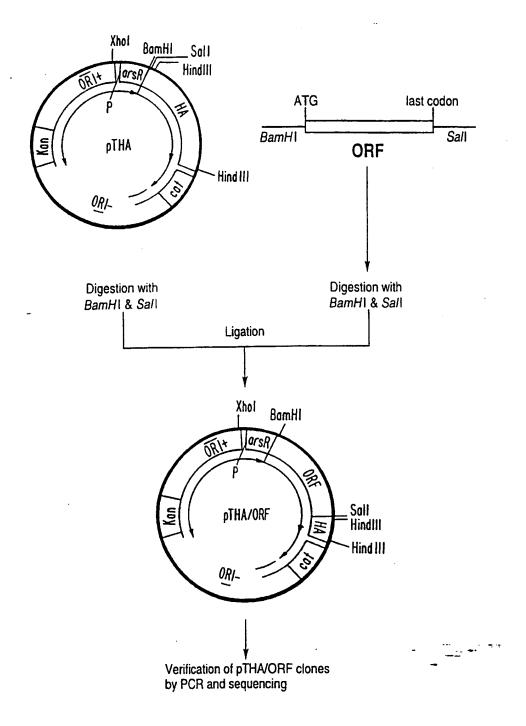


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FIG. 2.



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## FIG. 3

### (A) Functional assay on semi-solid support media

Frozen stock of phage 77 pTHA/ORF *S. aureus* RN4220 transformants

1:10 and 1:100 dilution in saline solution

5 µl of 1:10 dilution

3 µl of 1:10 and 1:100 dilution

Streak onto agar plates containing
0, 2.5, 5, and 7.5 µM NaAsO2

O/N, 37°C

Compare bacterial growth on plates with and without NaAsO2

Functional assay in liquid medium

# O/N culture inoculated from frozen stock of phage 77 pTHA/ORF *S. aureus* RN4220 transformants 1:100 dilution of O/N culture 2 h, 37°C, 250 rpm Fresh culture 150 µl 2.5 ml containing 0 and 5 µM NaAsO<sub>2</sub> 3.5 h, 37°C, 250 rpm Measure OD<sub>565</sub> 1:10 serial dilution from 10<sup>-1</sup> to 10<sup>-6</sup>

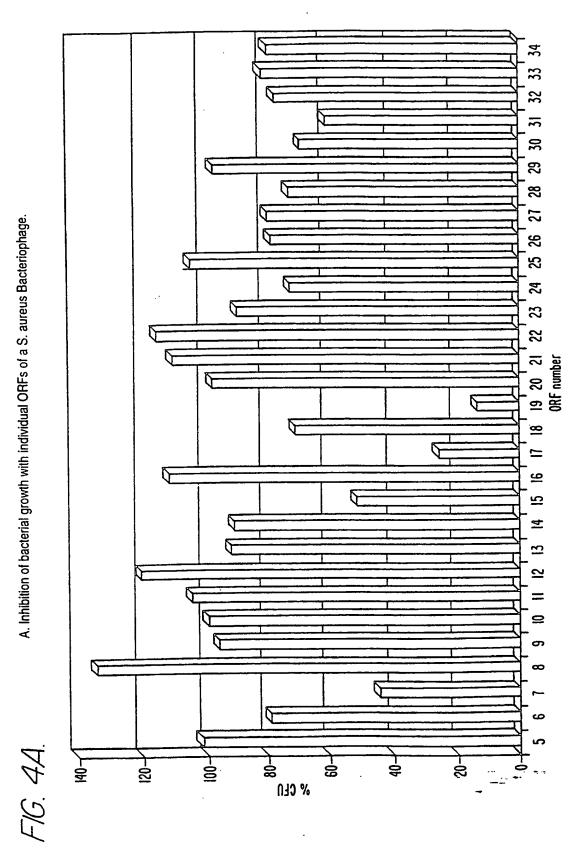
20 μl of 10<sup>-4</sup> to 10<sup>-6</sup>.

Spot onto agar plate

Count colonies

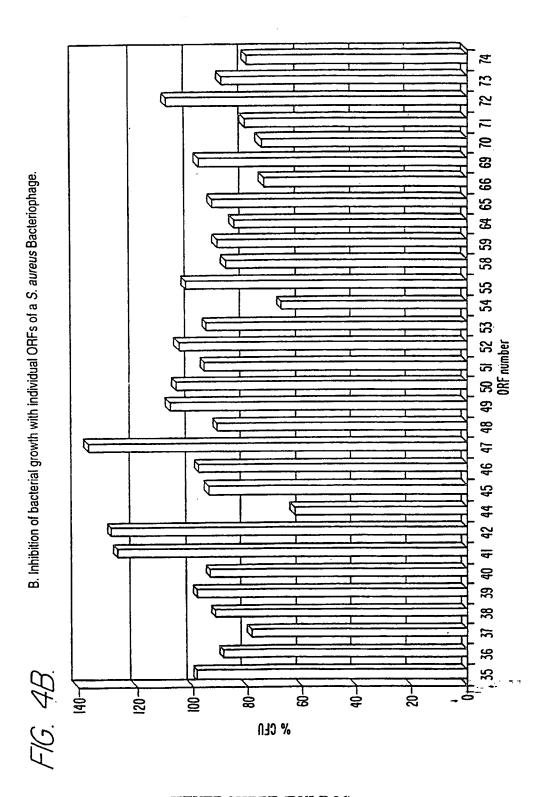
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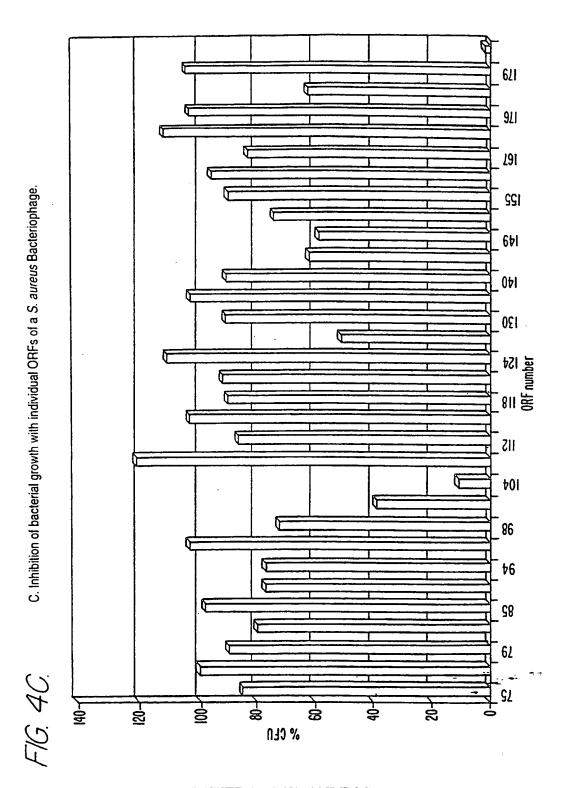
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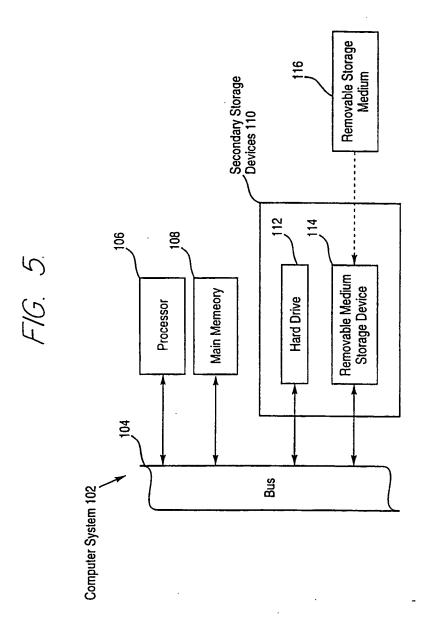
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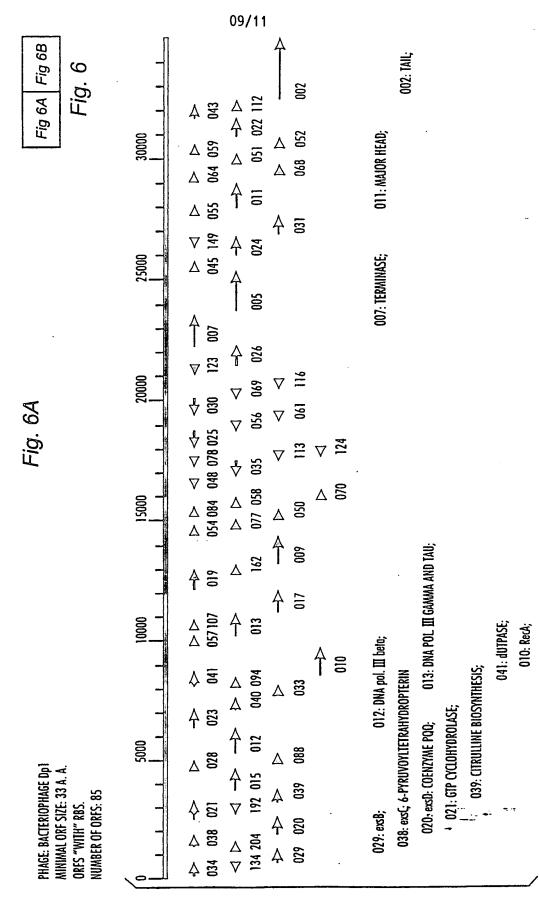
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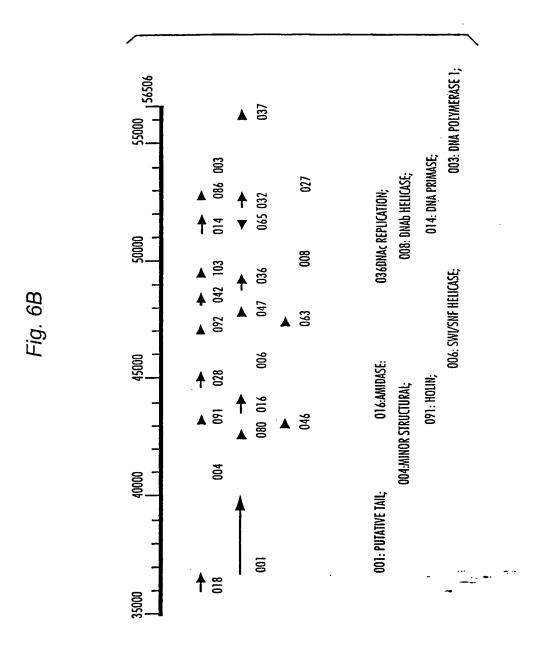
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# FIG. 7.

### Abbreviations:

kan: gene encoding kanamycin resistance
cat: gene encoding chloramphenicol resistance
ori + and -: origin of replication in gram-positive and
gram-negative bacteria, respectively
arsR: gene encoding regulatory protein of the ars promoter
P: ars promoter
lucFF: gene encoding luciferase protein. This portion will
be removed and replaced by individual *S. aureus* phage
genes.

### Referance:

Tauriainen et al., Appl. Environ. Microbio. 1997. 63: 4456-4461

